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Reliv International 2010 Annual Report

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Shareholder Information

Reliv International, Inc. is a developer, manufacturer and marketer of a proprietary line of nutritional supplements addressing basic nutrition, specific wellness needs, weight management and sports nutrition. Reliv also offers a line of premium skin care and food products under its Relivables brand. These products are sold exclusively to customers through Independent Reliv Distributors working in fifteen countries: United States, Australia, New Zealand, Canada, Mexico, United Kingdom, Ireland, the Philippines, Malaysia, Singapore, Germany, Austria, the Netherlands, Brunei and Indonesia.

2010 Financial Highlights

(In thousands, except per share amounts)

At December 31	2010	% change	2009
Net sales	\$ 78,748	(7.8)	\$ 85,399
Net income	1,683	(33.1)	2,515
Earnings per share			
Basic	0.14	(30.0)	0.20
Diluted	0.14	(30.0)	0.20
Total assets	24,844	2.9	24,154
Long-term debt, less current maturities	4,151	(12.1)	4,720
Stockholders' equity	13,931	13.6	12,267
Return on net sales	2.1%		2.9%
Return on average total assets	6.6%		10.1%
Return on equity	12.8%		18.6%
Current ratio	2.07		1.81



Dear Fellow Reliv Shareholder,

As we start 2011, I believe Reliv has the strategies in place to resume our strong sales growth. A great new product we introduced in the first quarter of 2011 has the potential to help boost sales, to attract new distributors, and to reenergize our brand.

We are well positioned and fundamentally sound. In a society that values health and financial stability, Reliv offers both, through nutritional supplements and business opportunities.

In short, I feel good about our 2011 prospects, which are discussed in detail below. First, I'll review Reliv's 2010 results.

2010 Results

Our financial results in 2010 were disappointing. Net sales of \$78.7 million represented a decline of 7.8 percent from 2009 net sales. We reported net income of \$1.7 million down 33.1 percent from 2009. Earnings per diluted share were \$0.14 compared to \$0.20 per diluted share in 2009.

International net sales were strong throughout the year, ending with an increase of 14.4 percent above 2009 international net sales. Our international markets continue to represent great growth potential for Reliv and we remain intent on capitalizing on the opportunity.

Net sales in the United States dropped 10.9 percent to \$66.9 million. We must generate sales growth in the U.S., and I'll discuss ways we intend to achieve that growth in the next section of this letter.

Despite a growing economy, poor job growth still appears to be dampening consumer spending among our distributors and customers.

Reliv remains in solid financial shape. We ended the year with cash and cash equivalents of \$6.3 million. We generated cash from operating activities of \$2.2 million during 2010.

We lowered our long-term debt by 10 percent during the year, and our long-term debt-to-equity is a low 0.34 to 1.



Energy Boost

To increase sales in 2011, we must:

- Sponsor new distributors into Reliv and attract new retail customers
- Maintain our already high retention rate
- Increase the size of our average order

We believe that Reliv 24K™, our first new nutritional product in three years, can contribute to each of those goals. Reliv 24K, an energy shot beverage, provides healthy energy and greater mental focus, while reducing stress.

It represents two firsts for Reliv. One: 24K is Reliv's first bottled drink. Two: 24K is Reliv's first product in the rapidly growing energy drink market. Here's why we think 24K will be a winner for Reliv.

With annual sales of \$7.6 billion, energy drinks are the fastest-growing market in the functional food industry. Within that market, the energy shot category accounts for \$560 million in annual sales. We seek to capitalize on the category by offering 24K in 28-ounce plastic bottles and in smaller double-shot bottles for on-the-go convenience.

Because it enhances mental focus, 24K also fits into the market for brain and memory supplements, another growing segment of the functional food industry. Annual sales in 2009 totaled \$385 million, twice as much as reported in each of the previous two years.

Both energy drinks and mental-clarity products particularly appeal to the Millennial Generation, which is now 18 to 29. We expect distributors to use 24K to introduce more Millennials to the Reliv family of products.

The unique nature of 24K, however, makes it ideal for everyone who leads an active life. It offers healthy energy and greater mental focus while reducing stress, which will appeal to consumers in all age groups. We also believe the ready-to-drink formulation will attract consumers who haven't yet experienced Reliv products.

We believe that many existing distributors and customers will add 24K to their standard order, which may lead to an increase in average order sizes. If the overall success of 24K meets our expectations, our research and development staff may pursue more ready-to-drink products.

The initial response to 24K has been terrific. The introduction at Reliv's 2011 U.S. National Conference in early February electrified the distributors in attendance, and the excitement has spread throughout the U.S. field. We think 24K will provide an energy boost for our sales.



Refreshing Reliv's Brand

In 2010, we began the long-term process of refreshing Reliv's brand, with a particular focus on associating Reliv with a healthy, active lifestyle. The Team Reliv Program plays a major role in that effort. The program consists of a series of running events sponsored by Reliv and its distributors throughout the United States and in our international markets. Reliv offices in Mexico and in the Philippines, for example, actively sponsor running and other athletic events.

Such events help distributors find prospective new customers and distributors in a positive environment. Distributors usually set up booths with product literature and free samples of our Innergize!® isotonic sports drink. This year, 24K samples may be offered too.

Not all of our customers find us at athletic events, though. Some of them find us in cyberspace. We have expanded our brand widely into social media and we plan to continue to expand our footprint in that area in 2011. We now have a thriving Facebook community of about 7,500 people, a number that continues to grow. Team Reliv also has its own Facebook page.

We continue to attract followers on Twitter. We have also established a Reliv YouTube channel as well as a corporate blog for distributors. All of these social media initiatives enhance the Reliv community, strengthen our relationships with distributors in the field, and reinforce our retention efforts.

Increase Sponsoring

We will continue our efforts to increase sponsoring. One of our goals for 2011 is to help our distributors welcome more new people into Reliv. Distributors who are already successful sponsors may be offered incentives to bring even more people into Reliv.

While focusing on growth opportunities, management has also been keeping costs under control. We continually improve our technology infrastructure and our offerings so that our distributors can operate more efficiently. We also constantly look for ways to enhance our internal efficiency. The goal, as sales turn around, is to ensure that a large percentage of the sales increase flows to the bottom line.

With the new 24K healthy energy shot, we remain true to our long-standing product strategy to make nutrition simple. Our nutritional products promote optimal health. Our business opportunities offer individuals potential financial stability and personal growth. And our charitable foundation combats hunger around the world. (See page 6 for more on the Reliv Kalogris Foundation.)

I want to thank our shareholders for their continued support. I also offer thanks to the finest and most dedicated distributors in the business. Finally, thank you to our hard-working staff members, who are always committed to excellence. I look forward to reporting on our progress to all of you next year.



Robert L. Montgomery
Chairman, President and Chief Executive Officer





Reliv Kalogris Foundation

2010 was a great year for the Reliv Kalogris Foundation, the charitable arm of Reliv International. Our generous distributors donated a record \$1.2 million to the foundation in 2010, enabling the organization to provide free nutritional supplements to malnourished people around the world.

Today, we donate supplements to feeding centers in 10 countries. We feed more than 42,000 people every day. Since 1995, the Foundation has donated \$26 million in nutritional products to programs that feed the hungry.

Highlights for 2010 include:

- Donation of funds to build a feeding center in the Philippines
- Donation of money to break ground for an orphanage in Haiti
- Raising more than \$40,000 at our fourth annual "Walk for the Mission" last summer in St. Louis
- A special donation of \$450,000 in Reliv nutritional supplements to victims of the Haiti earthquake in January 2010
- Establishment of the Network to Nourish Program, now registered in 48 states, which enables the organization to accept donations from any individual

The Reliv Kalogris Foundation spends almost 95 percent of all donations received directly on program services. That is well above the 75 percent criterion established by the American Institute of Philanthropy for its "Top Rated" list.

For more information on the Foundation, please visit relivkalogrisfoundation.org. Or keep up with us at facebook.com/RelivKalogrisFoundation.

Corporate Officers

Robert L. Montgomery
Chairman, President and
Chief Executive Officer

Carl W. Hastings, Ph.D.
Vice Chairman
Chief Scientific Officer

R. Scott Montgomery
Executive Vice President,
Chief Operating Officer

Ryan A. Montgomery
Executive Vice President,
Worldwide Sales

Steven D. Albright
Senior Vice President, Finance
Chief Financial Officer

Steven G. Hastings
Senior Vice President,
North American Sales

Stephen M. Merrick
Senior Vice President,
General Counsel and Secretary

Donald E. Gibbons, Jr.
Senior Vice President

Brett M. Hastings
Vice President, Legal

Debra P. Hellweg
Vice President, Operations

Ronald W. McCain
Vice President, Sales Development

Barry A. Murov
Vice President, Corporate Communications

Joseph J. Wojcik
Vice President, International

Kurt C. Wulff
Vice President, Marketing

Board of Directors

Robert L. Montgomery
Chairman, President and
Chief Executive Officer
Reliv International, Inc.

Carl W. Hastings, Ph.D.
Vice Chairman
Reliv International, Inc.

Stephen M. Merrick
Senior Vice President,
Reliv International, Inc.

Donald L. McCain
Corporate Secretary,
The Baughan Group, Inc.

John B. Akin
Retired Vice President,
A. G. Edwards, Inc.

Robert M. Henry
Executive Chairman
Immunotec Inc.

Denis St. John, CPA
Chairman
Real Estate Development Strategies, LLC

Michael D. Smith
Private Investor

John M. Klimek
Managing Director
HFR Asset Management

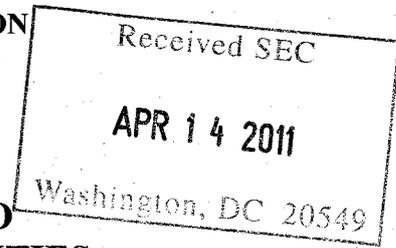
Five-Year Financial Summary

<i>(In thousands, except per share amounts)</i>	2010	2009	2008	2007	2006
Net sales	\$ 78,748	\$ 85,399	\$ 98,195	\$111,058	\$117,467
Net income	1,683	2,515	2,881	5,041	7,898
Earnings per common share:					
Basic	0.14	0.20	0.19	0.31	0.48
Diluted	0.14	0.20	0.19	0.31	0.47
Cash dividends per share of common stock	0.04	0.07	0.10	0.10	0.10
Total assets	24,844	24,154	23,893	33,607	37,282
Long-term debt, less current maturities	4,151	4,720	—	—	—

Stock Price & Dividend Summary

2010	High	Low	Close	Dividend
First Quarter	\$ 3.48	\$ 2.87	\$ 2.90	\$ —
Second Quarter	3.10	2.25	2.42	0.02
Third Quarter	2.43	2.00	2.15	—
Fourth Quarter	2.18	1.64	1.94	0.02
2009	High	Low	Close	Dividend
First Quarter	\$ 5.04	\$ 3.40	\$ 3.49	\$ —
Second Quarter	4.80	2.08	3.42	0.05
Third Quarter	4.12	2.85	3.34	—
Fourth Quarter	3.75	2.97	3.28	0.02

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, D.C. 20549



FORM 10-K
ANNUAL REPORT PURSUANT TO
SECTION 13 OR 15(d) OF THE SECURITIES
EXCHANGE ACT OF 1934

For the Fiscal Year Ended December 31, 2010

(Mark One)

ANNUAL REPORT PURSUANT TO SECTION 13 OR 15 (d) OF THE SECURITIES
EXCHANGE ACT OF 1934

For the fiscal year ended December 31, 2010

OR

TRANSITION REPORT PURSUANT TO SECTION 13 OR 15 (d) OF THE SECURITIES
EXCHANGE ACT OF 1934

For the transition period from _____ to _____

Commission File Number
000-19932

RELIV' INTERNATIONAL, INC.

(Exact name of Registrant as specified in its charter)

Delaware

(State or other jurisdiction of
incorporation or organization)

371172197

(I.R.S. Employer Identification Number)

136 Chesterfield Industrial Boulevard
Chesterfield, Missouri
(Address of principal executive offices)

63005
(Zip Code)

(636) 537-9715

Registrant's telephone number, including area code

Securities registered pursuant to Sections 12(b) of the Act:

Title of Each Class

Name of Each Exchange on Which Registered

Common Stock, par value \$0.001

NASDAQ Global Select Market

Securities registered pursuant to Section 12(g) of the Act: None

Indicate by check mark if the registrant is a well-known seasoned issuer, as defined in Rule 405 of the Securities Act. Yes No

Indicate by check mark if the registrant is not required to file reports pursuant to Section 13 or Section 15(d) of the Act. Yes No

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the

registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. Yes No

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Web site, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§ 232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit and post such files). Yes No

Indicate by check mark if disclosure of delinquent filers pursuant to Item 405 of Regulation S-K (§229.405 of this chapter) is not contained herein, and will not be contained, to the best of registrant's knowledge, in definitive proxy or information statements incorporated by reference in Part III of this Form 10-K or any amendment to this Form 10-K.

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, or a non-accelerated filer. See definition of "accelerated filer and large accelerated filer" in Rule 12b-2 of the Exchange Act. (Check one):

Large accelerated filer Accelerated filer Non-accelerated filer Smaller Reporting Company

Indicate by check mark whether registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). Yes No

Based upon the closing price of \$2.42 per share of the registrant's common stock as reported on the NASDAQ Global Select Market on June 30, 2010, the aggregate market value of the common stock held by non-affiliates of the registrant was approximately \$17.9 million. (The determination of stock ownership by non-affiliates was made solely for the purpose of responding to the requirements of the Form and the registrant is not bound by this determination for any other purpose.)

The number of shares outstanding of the registrant's common stock as of March 1, 2011 was 12,450,808 (excluding treasury shares).

DOCUMENTS INCORPORATED BY REFERENCE

<u>Document</u>	<u>Part of Form 10-K into Which Document Is Incorporated</u>
Sections of the registrant's definitive Proxy Statement for the Annual Meeting of Stockholders to be held on May 26, 2011, which is expected to be filed no later than 120 days after December 31, 2010	Part III

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FORWARD-LOOKING STATEMENTS

This annual report includes both historical and “forward-looking statements” within the meaning of Section 21E of the Securities Exchange Act of 1934, as amended. We have based these forward-looking statements on our current expectations and projections about future results. Words such as “may,” “should,” “could,” “would,” “expect,” “plan,” “anticipate,” “believe,” “estimate,” “predict,” “potential,” “continue,” or similar words are intended to identify forward-looking statements, although not all forward-looking statements contain these words. Although we believe that our opinions and expectations reflected in the forward-looking statements are reasonable, we cannot guarantee future results, levels of activity, performance or achievements, and our actual results may differ substantially from the views and expectations set forth in this annual report. We disclaim any intent or obligation to update any forward-looking statements after the date of this annual report to conform such statements to actual results or to changes in our opinions or expectations.

PART I

Item No. 1 - Business

Overview

We are a developer, manufacturer and marketer of a proprietary line of nutritional supplements addressing basic nutrition, specific wellness needs, weight management and sports nutrition. All but two of our science-based supplements are packaged in powdered form and are not only simple to use but also, when mixed with water, juice or other liquid and consumed, provide an effective means of delivering nutrients to the body. We also offer a ready-to-drink product, an encapsulated product and a line of skin care and food products. We sell our products through an international network marketing system using independent distributors. We have sold products in the United States since 1988 and in selected international markets since 1991.

We currently offer 17 nutritional supplements. In addition, we market a line of 15 skin care and food products under our Relivables brand. We have selectively evolved our product offering over our history. Our core line of nutritional supplements, which represented 63.1% of net product sales for the year ended December 31, 2010, includes the following four products:

- Reliv Classic and Reliv NOW — two basic nutritional supplements containing a full and balanced blend of vitamins, minerals, proteins and herbs
- Innergize! — an isotonic sports supplement in three flavors
- FibRestore — a high-fiber and antioxidant supplement

These are our most successful supplements based on fiscal year 2010 net sales. We have 13 other nutritional supplements that complement these four core products. We periodically refine our products and introduce related new products and product categories. Our internal research and development team has developed most of our products, and we hold U.S. patents on five of these products — Innergize!, FibRestore, Arthraffect, ReversAge and Cellebrate. In addition, we have applied for U.S. patents on our ProVantage, GlucAffect and CardioSentials products.

We believe that our network marketing model is the best method for the marketing and sale of our products because it utilizes ongoing personal contact among our distributors and their retail customers. This enables our distributors to communicate directly regarding the products, the business opportunity we offer and their personal experiences with both. We provide our distributors with a financially rewarding and entrepreneurial opportunity, affording them the ability to earn compensation both from the direct sale of products and from sales volume generated by distributors they sponsor. We actively support our distributors by providing marketing materials, a dependable product fulfillment system and frequent educational, training and motivational programs.

The majority of our sales traditionally has been, and is expected to continue to be, made through our distributors in the United States. We also currently generate sales through distributor networks in Australia, Austria, Brunei, Canada, Germany, Indonesia, Ireland, Malaysia, Mexico, the Netherlands, New Zealand, the Philippines, Singapore and the United Kingdom. In each country in which we conduct business, our distributors operate under a uniform business and compensation model that maintains consistent marketing, sales, fulfillment, and compliance

procedures. As of December 31, 2010, our network consisted of approximately 60,740 distributors — 47,450 in the United States and 13,290 across our international markets.

We manufacture all of our powdered nutritional supplements at our facility in Chesterfield, Missouri. We believe our ability to formulate and manufacture all but two of our own nutritional supplements enables us to produce our products efficiently while maintaining our high standards of quality assurance and proprietary product composition.

Industry Overview

Nutritional Supplement Market

We operate primarily in the \$26.9 billion U.S. nutritional supplement market, which is part of the broader \$102 billion U.S. nutrition industry according to 2009 data published by the *Nutrition Business Journal*, or NBJ, and \$228.0 billion global nutrition industry, also according to the NBJ, with sales expected to increase six to ten percent.

A combination of demographic, healthcare and lifestyle trends are expected to drive continued growth in the nutritional supplement market. These trends include:

- *Aging Population:* The older population (persons 65 years or older) numbered 39.6 million in 2009 according to the Department of Health and Human Services. They represented 12.9% of the U.S. population, about one in every eight Americans. By 2030, there will be about 72.1 million older persons living in the United States, more than twice their number in 2000. People 65 years or older represented 12.4% of the population in the year 2000 and are expected to represent 19% of the population by 2030. We believe this ever-growing population will continue to focus on their nutritional needs as they age.
- *Rising Healthcare Costs and Use of Preventative Maintenance:* The costs of healthcare in the United States continue to increase rapidly each year. National health care spending reached \$2.6 trillion in 2010 and is expected to reach \$4.5 trillion by 2019 according to the National Coalition on Health Care, or NCHC. In an article published by Families USA, they reported that nearly 6.9 million Americans lost their health coverage between 2008 and 2009. In addition, the total 2010 medical costs for a typical American family of four topped out at \$18,074 which is a 7.8% increase from 2009 according to the Milliman Medical Index. This is the highest dollar increase since the inception of the Milliman Medical Index 10 years ago. In order to maintain quality of life as well as reduce medical costs, many consumers take preventative measures to improve their general health, including the use of nutritional supplements.
- *Increasing Focus on Weight Management:* According to a report published in the January 2009 *Obesity* analyzing NHANES (The National Health and Nutrition Examination Survey) data, 86.3% of Americans will be overweight or obese by 2030, and less than 1% will be of normal weight by 2048 if the current weight trends continue. Related health care costs to obesity are expected to grow between \$860.7 billion to \$956.9 billion by 2030 and account for 16% to 18% of all medical expenditures. Being overweight can lead to more serious health concerns such as diabetes, heart disease and other chronic illnesses and individuals who are obese have a 10% to 50% increased risk of death from all causes, compared with healthy weight individuals. Bearing these facts in mind, we believe that there will be an increased need not only for weight loss products but for wellness products as well.

Direct Selling Market

Health and nutrition products are distributed through various market means, including retailers such as supermarkets, drugstores, mass merchants and specialty retailers; direct marketers such as mail order companies and Internet retailers; and direct sellers such as network marketers and healthcare practitioners. We distribute our products through the direct selling channel via our network marketers.

Direct selling involves the marketing of products and services directly to consumers in a person-to-person manner. Direct selling is a significant global industry largely utilized for the sale of a wide range of consumer products from companies such as Avon Products Inc., Alticor Inc. (Amway Corp.) and Tupperware Brands Corporation. According to the World Federation of Direct Selling Associations, or WFDSA, the 2009 global direct selling market (for all product categories) was estimated to be \$117.5 billion. The WFDSA estimates that the number of individuals engaged in direct selling nearly doubled between 1999 and 2009, from 35.9 million sellers to 74 million in 2009. The U.S. had 16.1 million direct sellers in 2009.

While the United States is currently the largest direct selling market with \$28.3 billion in annual sales in 2009, international markets account for 76% of the entire industry, according to the WFDSA. Eighteen countries (including the United States) have annual direct sales revenue of at least \$1 billion and another 26 countries have annual direct sales revenue of at least \$100 million, according to the WFDSA.

For the nutrition industry, the network marketing channel accounted for approximately 7% of the total U.S. nutritional supplements sold in 2008, or approximately \$7.1 billion, according to the NBJ.

We believe that we are well positioned to capitalize on the domestic and international growth trends in direct sales, as both a developer and manufacturer of proprietary nutritional products, utilizing our network marketing distribution system.

Our Competitive Strengths

We believe that we possess a number of competitive strengths that have historically supported our growth and enabled us to achieve sustained profitability.

Complete, Simple Nutrition. We focus on the completeness, balance and simplicity of our basic nutritional supplements — Reliv Classic and Reliv NOW — as captured by our slogan, “Nutrition Made Simple. Life Made Rich.” Because these two basic nutritional supplements each contain a full and balanced blend of vitamins, minerals, proteins and herbs, supplementation is made simple for the consumer, who does not have to select and purchase several supplements for his or her basic nutritional needs. For more specific individual needs, we provide 15 additional supplements. We believe that our two basic nutritional supplements, together with our additional supplements and Relivables products, enhance the ability of our distributors to build their businesses by providing a comprehensive, simple product offering.

Nutritional Supplements Consumed in Liquid Form. We believe that our nutritional supplements which are consumed in liquid form provide a competitive advantage over other supplements such as vitamins, minerals and herbs in pill or tablet form. Our powder-based nutritional products are consumed with water, milk or juice and 24K, our newest product, is a ready-to-drink product. Our products provide an effective means of delivering nutrients to the body. We believe nutrients taken orally in liquid form lead to better absorption at the cellular level, or “bioavailability.”

In-House Development and Production. We have developed substantially all of our nutritional supplement and food products utilizing nutrition science as the basis for product formulation. We maintain an ongoing research and development effort led by Carl W. Hastings, Ph.D., our Chief Scientific Officer and Vice Chairman. In addition, we consult regularly with other industry professionals and with the physicians on our Medical Advisory Board with respect to developments in nutritional science, product enhancements and new products. Since 1993, we have manufactured substantially all of our nutritional products at our facility in Chesterfield, Missouri. Currently, we outsource two nutritional supplement products, our Slimplicity accelerator capsules and 24K. We believe our ability to formulate and manufacture all but two of our own nutritional supplement products enables us to maintain our high standards of quality assurance and proprietary product composition.

Experienced Ambassador Team. Our Ambassador corps consists of distributors who have achieved the level of Master Director, have earned royalty payments of at least \$4,000 in consecutive months and meet our leadership and character criteria necessary to garner our invitation to be an Ambassador. Our Ambassadors generally are our most productive distributors and are essential in recruiting, motivating and training our entire distributor network. We, and our Ambassadors, lead hundreds of annual events throughout all of our markets to motivate and train distributors, including regular recruiting meetings, trainings, conference calls, training schools for Master

Affiliates and higher levels and regional, national and international distributor conferences. As of December 31, 2010, we had a total of 371 Ambassadors. The top 10 distributors at the Ambassador level have been with us for an average of 18 years, which provides consistency in training new distributors and contributes to increased sales.

Uniform Distributor Business Model. Our distributor compensation system is essentially uniform throughout our domestic and international markets. The compensation plan is “seamless” in that distributors in each market all receive discounts and commissions on relatively the same terms, subject to a few variances to address market conditions and cultural preferences. We also provide consistent distributor documentation and training throughout our system and in all of our markets. We believe this uniform model is effective in motivating and training distributors to build their businesses and enter new markets.

Experienced and Incentivized Management Team. Our management team is led by our founder, Robert L. Montgomery, who has been our Chief Executive Officer since the inception of our company in 1985. Our executive officers have been employed by our company for an average of 16 years and are experienced in their areas of focus, which include manufacturing, sales, finance, marketing and operations. As of March 1, 2011, our directors and executive officers beneficially own approximately 40.7% of our common stock.

Our Business Strategy

Our basic objective is to increase our net sales by increasing the number and productivity of our distributors and by periodically improving our existing products and introducing new products. We also intend to invest in our infrastructure to improve our operating efficiencies, provide better service to our distributors and leverage our current operating facilities to improve our profitability. We seek to accomplish these objectives by employing the following strategic initiatives:

Leverage and Expand our Existing Distributor Base Throughout the United States. The United States has been and will continue to be our largest market. Our growth strategy in the United States involves multiple initiatives, such as continued investment in company-sponsored events and training and better utilization of our upper-level distributors across different geographical areas.

Expand in Existing and New International Markets. We believe there is a significant opportunity to increase our net sales in international markets. We have a uniform business model across all of our markets and encourage our distributors to pursue their business in multiple markets. In selected markets, we have begun investing in additional marketing support for our distributors that is consistent with our successful activities in the United States, including third party advertising materials and company-sponsored distributor meetings. We believe this uniform business model and additional marketing expense will encourage expansion of our distributors in our existing international markets and will provide a framework that facilitates our entry into new international markets. To that end, we continue to monitor business conditions in potential new markets and will selectively expand as timing and conditions are appropriate.

Invest in Improved and New Products. As a developer of nutritional supplements, it is vital to continue to invest in the research and development of new and innovative products. Additionally, we will continue to improve and validate the efficacy of our existing product line. For example, in February 2011 we launched 24K, our first ready-to-drink product, to support energy production and mental focus and in August 2009 we introduced our Relivables brand of products. These types of investments should facilitate customer and distributor retention, as well as the recruitment of new distributors.

Expand and Improve our Manufacturing and Distribution Capabilities. We currently manufacture all of our powdered nutritional supplements at our facility in Chesterfield, Missouri. This allows us to precisely control product composition and quality assurance. Periodically, we make appropriate investments that enhance our manufacturing capabilities and capacity to further leverage our existing facilities and trained production staff. We expect to continue to make appropriate investments in our manufacturing and fulfillment facilities.

Increase Appeal to Broader Demographic. Traditionally, our customer and distributor demographic has skewed towards baby boomers and older individuals searching for nutritional solutions to supplement their diet and support overall wellness. While continuing to maintain our focus on the needs of this important segment, we believe there is an opportunity to expand our sales and distributor base by increasing our appeal to younger generations

interested in nutrition and an active healthy lifestyle. We believe the nutritional aspects and convenience of 24K, our healthy energy and mental focus drink, will attract health conscious on-the-go individuals, many of whom fall within the under-40 demographic. In addition, we initiated the Team Reliv program in February 2010 that provides financial and marketing support to local distributor groups sponsoring races and walks within their local communities. Further, we maintain an active presence on popular social media sites including Facebook, Twitter, YouTube and several other social networks that are popular with younger generations. Our internal social media team is comprised of Gen X and Gen Y staffers who regularly interact with distributors, customers and prospects. We plan to continue to develop products and programs, and expand our technology offerings in an effort to further appeal to younger generations interested in healthy active lifestyles and a vibrant evolving business opportunity.

Our Products

Product Overview

Our product line includes nutritional supplements that address basic nutrition, specific wellness needs, weight management and sports nutrition. We combine ingredients from science and nature in targeted, well-balanced, easy-to-use formulas that are specifically designed to enhance wellness and increase performance and energy in specific applications. All but two of our supplements are in powdered form that the consumer mixes with water, juice or other liquid. We also have a ready-to-drink product, an encapsulated product and a line of skin care and food products marketed under our Relivables brand name.

We currently offer 17 nutritional supplements. In addition, we offer 11 skin care and four food products under our Relivables line. Our basic nutritional supplements are formulated to provide a balanced and complete level of supplementation for the consumer. For more specific needs, we provide other focused product formulations. We have purposely been selective in the number and types of products that we offer. By providing a line of targeted products, we make it simple for our distributors and consumers to choose products appropriate for their objectives. We consider four of our oldest and best selling products — Reliv Classic, Reliv NOW, Innergize!, and FibRestore — to be our primary or “core” products.

The following table summarizes our product categories as of December 31, 2010. The net sales figures are for the year ended December 31, 2010:

<u>Product Category</u>	<u>Product Name</u>	<u>% of 2010 Net Sales⁽¹⁾</u>	<u>Year Introduced</u>
Basic Nutrition	Reliv Classic.....	18.1	1988
	Reliv NOW.....	11.6	1988
	NOW for Kids.....	3.8	2000
	Reliv Delight.....	0.2	2001
Specific Wellness	FibRestore.....	14.0	1993
	Arthaaffect.....	7.1	1996
	ReversAge.....	4.7	2000
	SoySentials.....	2.1	1998
	CardioSentials.....	2.2	2005
	GlucAffect.....	2.8	2008
Weight Management⁽²⁾	Simplicity Meal Replacement.....	1.9	2007
	Simplicity Accelerator Capsule.....	0.8	2007
	Reliv Ultrim Plus.....	0.3	1988
	Celebrate.....	0.7	1995
	Sports Nutrition	Innergize!.....	10.8
	ProVantage.....	3.3	1997
Relivables⁽³⁾	Skin Care.....	1.2	2001
	Food Products.....	0.7	2009

(footnotes on following page)

- (1) This table does not include net sales for the year ended December 31, 2010 related to freight and handling and sales of marketing materials, which represented approximately 13.7% of net sales for the year ended December 31, 2010.
- (2) Since its introduction in February 2007, our Slimplicity Meal Replacement formula has replaced Reliv Ultrim-Plus in all but our Canadian and Mexican markets. Upon introduction of our Slimplicity products in a particular market, our Reliv Ultrim-Plus line was discontinued in that market.
- (3) In August 2009, we introduced our Relivables line of skin care and food products. The Relivables skin care products include science-based updates to our original ReversAge skin care products. In conjunction with these updates, we have re-branded our skin care products under the Relivables lines.

Basic Nutrition Supplements

Our four basic nutrition supplements provide consumers with a broad spectrum of essential nutrients. Every formulation is specifically designed to optimize and enhance the benefits of the nutrients it contains.

- Reliv Classic is a nutritional supplement containing a variety of vitamins and minerals, soy and other protein sources and various herbs. It is a vegetarian product that contains no animal compounds, artificial preservatives, artificial flavors or added simple sugars. Reliv Classic is available in the United States, Australia, New Zealand, Canada, Germany, Austria, the Netherlands, the United Kingdom, Ireland, Malaysia, Singapore, Brunei and the Philippines.
- Reliv NOW is a nutritional supplement containing a variety of vitamins and minerals, soy and other protein sources and various herbs. Reliv NOW is available in every country where we operate except Indonesia.
- NOW for Kids is a product designed to provide a balanced nutritional supplement for a child's diet and contains a variety of vitamins and minerals. NOW for Kids is available in Australia, New Zealand, United States, the United Kingdom, Ireland, Austria, the Netherlands, Mexico, Malaysia, Brunei, and the Philippines.
- Reliv Delight is a powdered nutritional supplement marketed as a milk replacement. Reliv Delight is available in Mexico and the United States.

Specific Wellness Supplements

Our line of six specific wellness supplements contains specific compounds that target certain conditions and promote health. Each product is intended to work in conjunction with our basic nutritional supplement formulas to provide an effective, balanced and natural method for sustaining health and well-being.

- ReversAge is a patented youth-promoting nutritional supplement designed to slow down the effects of the aging process. Three proprietary complexes form the foundation of the supplement: longevity complex, antioxidant complex and herbal complex. The longevity complex is restorative and designed to replenish key hormones while creating balance within the body's major systems; the antioxidant complex is designed to slow aging at the cellular level and the herbal complex delivers a variety of herbs, including Ginkgo Biloba and Maca. ReversAge is available in every country where we operate except Germany, the United Kingdom, Ireland, and Indonesia. In Canada, the product is marketed as Nutriversal.
- SoySentials is a nutritional supplement containing soy as well as other vitamins, minerals and herbs designed for use by women. SoySentials provides a woman with key nutrients targeted to promote women's health and ease the symptoms of menopause and PMS. SoySentials is available in the United States and Mexico.

- CardioSentials is a berry-flavored nutritional supplement introduced in February 2005 that promotes heart health. The product contains 1,500 mg of phytosterols per serving, policosanol and several powerful antioxidants. In a clinical study of this product, participants experienced meaningful reductions in cholesterol as well as improvement in their high-density lipoprotein, or HDL, and low-density lipoprotein, or LDL, ratios. We have applied for a U.S. patent on CardioSentials. CardioSentials is available only in the United States.
- Arthraffect is a patented nutritional supplement containing Arthred, a patented form of hydrolyzed collagen protein, which is clinically reported to support healthy joint function. The product is available in the United States, Australia, New Zealand, Mexico, the Philippines, Malaysia, Singapore, Brunei and Canada. The product is marketed as A-Affect in Australia, New Zealand and Canada due to local product regulations.
- FibRestore is a patented nutritional supplement containing fiber, vitamins, minerals and herbs. A modified version of the FibRestore formula is marketed in Canada under the name Herbal Harmony to comply with Canada's nutritional regulations. FibRestore is available in all of the countries in which we operate except Indonesia.
- GlucAffect is a cinnamon cream flavored nutritional supplement launched in November 2008. GlucAffect contains Pycnogenol® and other clinically supported active ingredients. GlucAffect has been clinically proven to assist in healthy blood sugar management and support weight loss. We have applied for a U.S. patent on GlucAffect. GlucAffect is available in the United States, Canada, the Philippines, Malaysia, Singapore, and Brunei.
- 24K is our newest product, introduced in February 2011. 24K is our first ready-to-drink nutritional supplement available in a multi-serving 28-ounce bottle and in a two-ounce single serving bottle. 24K is formulated with a synergistic blend of 24 active ingredients designed to enhance the body's natural vitality and provide energy, focus and stress relief. It contains no caffeine and only 5 calories per serving. 24K is available only in the United States.

Weight Management Supplements

Our four weight management supplements combine advanced weight loss promoting complexes with scientifically balanced nutrition and health enhancing soy protein. Our ingredients are designed to work together, along with proper diet and exercise, to turn unwanted fat into energy without sacrificing muscle mass.

- Our Slimplicity Weight Loss System was introduced in the United States in February 2007 and includes two products: (1) Slimplicity meal replacement and (2) Slimplicity accelerator capsules. Our Slimplicity Weight Loss System incorporates these new products into an overall program that includes proper diet and exercise and is focused on facilitating weight loss and developing healthier lifestyle choices. Slimplicity is currently available in the United States, Germany, Austria, the Netherlands, Ireland, the United Kingdom, Australia, New Zealand, the Philippines, Malaysia, Singapore and Brunei. In Australia and New Zealand, the products are marketed as Slimsimply due to trademark availability.
- Reliv Ultrim-Plus is designed as a meal replacement (for a maximum of two meals per day) for use in a weight loss program. Reliv Ultrim-Plus is only sold in Canada and Mexico. Reliv Ultrim-Plus is no longer available in our other markets due to the introduction of our Slimplicity meal replacement product. We expect Slimplicity to eventually replace Reliv Ultrim-Plus in all of our markets.
- Cellebrate is a patented weight loss aid designed to suppress appetite, curb the storage of body fat, and facilitate the body's fat burning process. Cellebrate is available in the United States and Canada.

Sports Nutrition Supplements

Our two sports nutrition supplements contain a balance of nutrients scientifically designed to improve athletic performance and endurance, as well as muscle recovery and repair.

- Innergize! is a patented sports supplement, containing vitamins and minerals designed for performance enhancement. Innergize! is available in every country where we operate. In Canada, the product is marketed as Optain due to local product regulations.
- ProVantage is a nutritional supplement containing soy designed to enhance athletic performance with a balance of nutrients needed to improve endurance, muscle recovery and repair. ProVantage is designed to increase muscle recovery, muscle mass and function, reduce fatigue and burn excess body fat for extra energy. The product also benefits dieters and others seeking to increase their soy intake. We have applied for a U.S. patent on ProVantage. ProVantage is available in the United States and Canada.

Relivables

Our new Relivables product line is comprised of nutritionally sound skin care and food products. The new skin care products, marketed as the “r” skin care collection, are designed to create healthier, more youthful looking skin. Each product in our r collection contains the exclusive RA7 complex, an array of antioxidants, anti-inflammatory and anti-aging nutrients. These nutrients work together to slow the aging process and improve the skin’s appearance. The men’s “r” collection includes a body wash, shave lotion and after shave moisturizer with SPF 15. The women’s collection includes a cleansing facial wash, eye cream, daytime facial moisturizer with SPF 15, a nighttime facial moisturizer, and a body lotion. The r products are available in the United States, Australia and New Zealand.

The food products include Relivables All-Natural Sweetener, to be used in place of sugar or other artificial sweeteners. Its all-natural sweetener, derived from the stevia plant, has no sugar and contains one gram of fiber. Relivables Fortified Soy Milk is lactose-free and dairy-free and significantly exceeds the amount of calcium and vitamin D in dairy milk, along with six grams of soy protein. Relivables Soy Nuts are a good source of fiber and soy protein, low in sodium and cholesterol free. Relivables Healthy Snack Bars, which come in a Chocolate-Coated Granola flavor, are a good source of fiber, soy protein and whole grains.

Research and Development

We maintain an ongoing research and development effort, led by Carl W. Hastings, Ph.D., and consult with other industry professionals and with the physicians and professionals on our Medical Advisory Board with respect to developments in nutritional science, product enhancements and new products. Since 2000, we have introduced eight of our current nutritional supplement products, including ReversAge, NOW for Kids, Reliv Delight, GlucAffect, CardioSentials, Slimplicity meal replacement, Slimplicity accelerator capsules, and 24K. From time to time, we have also reformulated and enhanced our products. Our research and development team consistently evaluates product advancements in the marketplace and advancements in raw materials and ingredients available for new product ideas and developments.

For the years ended December 31, 2010 and 2009, our research and development expenses were \$587,000 and \$551,000, respectively.

Network Marketing Program

General Overview

We market and sell our products through a network marketing system of independent distributors, who purchase our products from us, or from other distributors, and who then sell our products directly to consumers. In addition to selling our products, our distributors also recruit others to distribute our products. Distributors receive compensation from both the sale of the products they have purchased at wholesale and, in the case of Master Affiliates and above, commissions on the volume of products sold by their downline organization. We believe network marketing is an effective way to distribute our products because it allows and relies on personal contact, education and endorsement of products which are not as readily available through other distribution channels.

We recognize that our sales growth is based on the continued development and growth of our independent distributor force and we strive to maintain an active and motivated distributor network through a combination of

quality products, discounts, commissions and bonus payments, sales conventions, training, personal recognition and a variety of publications and promotional materials.

Program Structure

Individuals who desire to market and sell our products may become distributors by being sponsored into the program by an existing distributor, and becoming part of that distributor’s “downline.” We offer a tiered discount and commission, or royalty, format that consists of four principal levels and several sub-levels, which are designed to compensate and motivate distributors to increase their networks and sales volumes.

Our distributors consist principally of individuals, although we also permit entities such as corporations, partnerships, limited liability companies and trusts to become distributors. A new distributor is required to complete a distributor application and, in most areas, to purchase a package of distributor materials (for \$25 plus shipping in the United States) consisting of a Distributor Guide and CD, business forms and promotional materials. The Distributor Agreement, when accepted by us, becomes the contract between us and the distributor and obligates the distributor to the terms of the agreement, which includes our Policies and Procedures for conduct of their business. All distributors are independent contractors and are not our employees.

In each country in which we conduct business, distributors operate under a uniform compensation system pursuant to which distributors generally are compensated based on their sales volumes. On the basis of sales volume or commission volume, distributors may achieve the following successive levels of achievement and compensation:

<u>Designation</u>	<u>Discount</u>
Retail Distributor.....	20%
Affiliate.....	25%
Key Affiliate.....	30%
Senior Affiliate.....	35%
Master Affiliate.....	40% ⁽¹⁾
Director.....	40% ⁽¹⁾
Key Director.....	40% ⁽¹⁾
Senior Director.....	40% ⁽¹⁾
Master Director.....	40% ⁽¹⁾
Presidential Director.....	40% ⁽¹⁾

⁽¹⁾ In addition to discounts, these levels also receive commissions based on sales in their downline organization.

Distributors purchase products from us at a discount from the suggested retail price for the products and then may sell the product at retail to customers, sell the product to other distributors at wholesale or consume the product. The amount of the discount varies depending on the distributor’s level of achievement, as indicated above.

Distributors generate income equal to the difference between the price at which they sell the product to customers and the discounted price they pay for the product. Distributors also earn wholesale commissions on products purchased by downline distributors in the distributor’s sponsored group equal to the difference between the price at which the distributor is entitled to purchase product and the price at which downline distributors purchase product. We calculate payments and issue a check directly to the qualified distributor once a month. For example, assume Distributor A is a 40% discount Master Affiliate who signs up Distributor B, a 30% discount Key Affiliate, who signs up Distributor C, a 20% discount Retail Distributor. If Distributor C purchases directly from us, a 10% wholesale profit check will be sent to Distributor A and B.

Upon achieving the level of Master Affiliate, distributors begin to receive additional compensation — “generation royalty” — payments of 8%, 6%, 4%, 3% and 2% of the retail volume of product purchased from us by Master Affiliates and above (and their personal groups) whom they have sponsored, and for each of five downline levels of sponsorship. To qualify for these additional compensation payments, Master Affiliates and above are required to maintain certain monthly sales volumes and to document specified levels of retail sales.

Master Affiliates who sponsor other distributors that achieve the level of Master Affiliate are entitled to become part of the Director Program. Advancement at the Director level is based upon achieving increasing levels of royalties based on sales generated by other distributors in the Director's downline organization. Distributors achieving each level receive recognition for their achievements at our company-sponsored events and in our publications. We also have a Star Director Program under which distributors achieving the level of Director and above receive additional compensation based on the number of Master Affiliates they have sponsored into the program. Directors receive an additional 1% to 3% royalty on the retail sales volume of Master Affiliates in their downline organization for an unlimited number of levels of sponsorship, until reaching a level that includes a Master Affiliate who also has achieved Star Director status.

Master Directors and Presidential Directors may also be invited to participate in the Ambassador Program. As of December 31, 2010, we had 371 Ambassadors. Qualifications to be invited by us to participate in the Ambassador Program include demonstrated competence and leadership qualities. Ambassadors receive recognition and awards for achieving Ambassador status and can then achieve additional levels of accomplishment. We utilize our Ambassadors to lead meetings and conferences, and to provide training and education to our distributors. Ambassadors achieving the level of Silver and higher also participate in the "Reliv Inner Circle," which may entitle them to receive additional compensation, paid participation in our sponsored events, health insurance and car allowances.

In addition to the levels of compensation described, we also provide a variety of incentives, bonuses, awards and trips to distributors who achieve high sales volumes and who advance in the distributor ranks.

Distributor Training, Motivation and Management

Our marketing efforts are focused on the development, training, motivation and support of our independent distributors. We support an active training program for our distributors in which our representatives and experienced distributors, usually Ambassadors, lead group training sessions. We provide distributors with manuals, brochures and other promotional, training and informational publications. We encourage distributors to hold regular Tuesday evening recruiting meetings and Saturday training sessions. We sponsor weekly training conference calls in which a significant number of distributors participate.

Our sponsorship generally includes the following:

- During 2010, we sponsored approximately 50 training schools on a quarterly basis across all of our markets for new Master Affiliates;
- For each market in which we operate, we sponsor an annual conference for distributors; and
- In the United States, we sponsor an annual International Conference in summer for all worldwide distributors and a winter conference for U.S. distributors.

During 2010, we invested approximately \$3.18 million in training, conferences and promotional events for our distributors worldwide.

Distributor Compliance

Our distributor organization and business model are designed and intended to promote the sale of our products to consumers by distributors. Sales training and promotional efforts emphasize that intention. To that end, we monitor purchases by distributors to identify potentially excessive individual purchases and keep detailed information regarding customer purchases through our corporate shopping cart and as part of our autoship program. Distributors are not required at any time to purchase product, although Master Affiliates and above are required to maintain certain minimum sales levels in their personal groups to continue receiving generation royalty compensation payments.

Distributors may create their own advertising provided that it is within our advertising rules. Unless a distributor is using our designed and approved advertisements, the distributor must submit for approval in writing all

advertising (e.g. brochures, flyers, audio tapes, classified or display ads, radio scripts) to our Compliance Department before placing it or arranging for placement.

Pursuant to our Policies and Procedures, which are incorporated by reference into our Distributor Agreement, distributors are permitted to make only those claims about our products that have been approved by us and/or provided in sales and training materials. Distributors acknowledge that our products are not represented as drugs and they are not authorized to make any diagnosis of any medical condition, make drug-type claims for, or prescribe our products to treat or cure, any disease or condition. We do not authorize or permit our distributors to make any express or implied references with regard to our products that they cure, prevent or relieve disease, replace or augment medication, provide therapy, promote healing, alleviate illnesses or symptoms of illnesses, or make any other medical claims for specific ailments.

In order to comply with regulations that apply to both us and our distributors, we conduct considerable research into the applicable regulatory framework prior to entering any new market to identify all necessary licenses and approvals and applicable limitations on operations in that market. We devote substantial resources to obtaining the necessary licenses and approvals and maintaining operations that are in compliance with the applicable limitations. We also research laws applicable to distributor operations and revise or alter distributor materials and products, as required by applicable regulations in each market.

Regulations in existing and new markets often are ambiguous and subject to considerable interpretive and enforcement discretion by the responsible regulators. In addition, regulations affecting our business often change and are subject to varying interpretation and application. We make every effort to monitor and comply with changes in laws and regulations as they occur.

We have a Compliance Department that receives and reviews allegations of distributor misconduct. If we determine that a distributor has violated our Policies and Procedures, we may take a number of disciplinary actions. For example, we may impose sanctions such as warnings or suspensions until specific conditions are satisfied, or take other appropriate actions at our discretion, including termination of the distributor's agreement.

Geographic Presence

Markets

We currently sell our products throughout the United States and in 14 other countries around the world. We have sold products in the United States since 1988 and sold our first product outside of the United States in 1991 when we entered Australia. In 2010, approximately 15.0% of our net sales were generated outside of the United States.

The table below shows the countries in which we operate and the year we commenced selling products:

<u>Country</u>	<u>Year Entered</u>	<u>Country</u>	<u>Year Entered</u>
United States	1988	Ireland	2003
Australia	1991	Singapore	2004
New Zealand	1992	Germany	2005
Canada	1992	Austria	2006
Mexico	1993	Netherlands	2006
United Kingdom ⁽¹⁾	1995	Brunei	2009
Philippines	2000	Indonesia	2009
Malaysia	2003		

⁽¹⁾ Includes Great Britain, Scotland, Wales and Northern Ireland.

Within the United States, we sell our products to distributors in all 50 states. We derived 35.5% of our net sales in 2010 in California, Illinois, Kansas, Texas, Missouri, Michigan, and Ohio, with each state contributing at least 4% of net sales. We believe that there is the opportunity to increase the number of our distributors in all markets where we sell our products, as our existing distributor bases grow and expand.

We organize all of our international operations under our wholly owned subsidiary, Reliv' World. As of December 31, 2010, Reliv' World consisted of the following market-specific entities: Reliv' Australia, Reliv' New Zealand, Reliv' Canada, Reliv' Mexico, Reliv' UK (including Ireland), Reliv' Philippines, Reliv' Malaysia, Reliv' Singapore, Reliv' Brunei, Reliv' Germany (including Austria and the Netherlands), and PT Reliv Indonesia. We have utilized this method of separate corporations in most of our markets, as local business licensing and product approvals require a local legal entity.

We believe that there is a significant opportunity to increase sales in all of our current international markets. We have established a uniform business model and compensation plan across all of our markets, and we continue to support our international markets with additional marketing programs and materials.

New Market Entry Process

We constantly evaluate new markets for our products. In order to do so, we perform an analysis of synergies between new and existing countries and distributor presence or interest in new markets, market conditions, regulatory conditions, product approval procedures and competition before selecting markets to enter. Once we decide to enter a new market, we first hire local legal counsel and/or a consultant with appropriate expertise to:

- help ensure that our network marketing system and products comply with all applicable regulations;
- help establish favorable public relations in the new market by acting as an intermediary between us and local regulatory authorities, public officials and business people; and
- explain our products and product ingredients to appropriate regulators and, when necessary, to arrange for local technicians to conduct required ingredient analysis tests of the products.

Where regulatory approval in a foreign market is required, local counsel and/or consultants work with regulatory agencies to confirm that all of the ingredients in our products are permissible within the new market. Where reformulation of one or more of our products is required, we attempt to obtain substitute or replacement ingredients. During the regulatory compliance process, we may alter the formulation, packaging, branding or labeling of our products to conform to applicable regulations as well as local variations in customs and consumer habits, and we may modify some aspects of our network marketing system as necessary to comply with applicable regulations.

Following completion of the regulatory compliance phase, we undertake the steps necessary to meet the operations requirements of the new market. In the majority of our new markets, we establish a sales center in a major city and provide for product purchases by telephone and/or pick up. Product is shipped to the purchaser from a warehouse located in the general geographic market or the distributor may walk in to the local office and purchase products, if a pick up center is available. In addition, we initiate plans to satisfy inventory, personnel and transportation requirements of the new market, and we modify our distributor materials, recordings, videos and other training materials as necessary to be suitable for the new market.

In some countries, regulations applicable to the activities of our distributors also may affect our business because in some countries we are, or regulators may assert that we are, responsible for our distributors' conduct. In these countries, regulators may request or require that we take steps to ensure that our distributors comply with local regulations.

Manufacturing

We established a manufacturing line at our headquarters facility in Chesterfield, Missouri and began to manufacture all of our nutritional supplements in early 1993. We expanded our Chesterfield facility in 1997 to now include 126,000 square feet of total space. At this facility, we manufacture all of our powdered nutritional supplements for distribution both domestically and internationally. Our Slimplicity accelerator capsules, 24K, and Relivables soy nuts and snack bars are manufactured by a third party and our Relivables skin care line is manufactured by third parties that are both owner and licensee of certain proprietary technology used in our skin care products.

Our ability to manufacture our powdered nutritional supplements is a competitive advantage over competitors not engaged in manufacturing and contributes to our ability to provide high-quality products. Our product manufacturing includes identifying suppliers of raw materials, acquiring the finest quality raw materials, blending exact amounts of raw materials into batches, and canning and labeling the finished products. Since we carefully select our ingredient suppliers, we are able to control the quality of raw materials and our finished products. We have not experienced any significant difficulty in obtaining supplies of raw materials for our nutritional supplements or finished product of our Slimplicity accelerator capsules or 24K. By monitoring and testing products at all stages of the manufacturing process, we precisely control product composition. In addition, we can control costs by manufacturing our own powdered nutritional supplements.

In 1996, we received approval from the Australian Therapeutic Goods Administration, or TGA, to manufacture products sold in Australia at our Chesterfield plant. The certification of our Chesterfield site by the Australian TGA also satisfied Canadian requirements. In 2010, our Chesterfield plant was audited and re-certified by the Australian TGA.

Fulfillment

Distributors order product in case lots of individual quantities and pay for the goods prior to shipment. We offer our Direct Advantage for distributors and their retail customers to order product in less than case lots directly from us by phone. Direct Advantage, an automatic monthly reorder program available for distributors and customers, provides a simple and convenient ordering process for consumers as well as distributors wanting to satisfy maintenance requirements. Product is shipped directly to the distributor or customer and upline distributors earn wholesale profits or, if applicable, a commission on all Direct Advantage sales.

In the United States, our products are warehoused at our Chesterfield facility and shipped by common carrier to distributors upon order. Our facility in Chesterfield, Missouri serves all parts of the country. Our products are also warehoused in, and shipped to local distributors from: Sydney, Australia; Auckland, New Zealand; Oakville, Canada; Birmingham, England; Petaling Jaya, Malaysia; Singapore; Brunei; and Jakarta, Indonesia. Our Philippines subsidiary currently has approximately three product pick-up centers located throughout the country which are operated by local business contractors and a company-owned and operated business center located in Makati. In Mexico, product is warehoused in and shipped from approximately five distribution centers located throughout the country. With the exception of our Canada, New Zealand, Singapore, Brunei, and German subsidiaries, each of our subsidiaries maintains an office and personnel to receive, record, and fill orders from distributors. Distributors in Ireland, Germany, Austria, and the Netherlands order and receive product from our UK subsidiary.

We maintain a policy that unused product may be returned by a customer to the selling distributor for a full refund or exchange within 30 days after purchase. We also maintain a policy that any distributor who terminates his or her distributorship may return saleable product which was purchased from us within twelve months of the termination for a refund of 90% of the purchase price less any compensation received relating to the purchase of the products. We believe this buyback policy addresses and satisfies a number of regulatory compliance issues pertaining to network marketing systems.

Historically, product returns and buy backs have not been significant. Product returns and buy backs have been approximately 0.67% and 0.70% of net sales in 2010 and 2009, respectively.

Information Technology Systems

In order to facilitate our continued growth and support distributor activities, we continually upgrade our management information and telecommunication systems, along with increasing our internet-based capabilities. These systems include: (1) a centralized host computer in our Chesterfield headquarters, which is linked to our international offices via secure frame relay connections that provide real-time order entry and information to respond to distributor inquiries, as well as financial and inventory management systems; (2) local area networks of personal computers within our markets, serving our local administrative staffs; (3) an international e-mail system through which our employees communicate; and (4) internet capabilities that provide a variety of online services to distributors, including product ordering, product information, event information and other related announcements,

and tools to assist distributor leaders in managing their downline distributor group. We continue to pursue initiatives to increase the percentage of distributor orders placed via the internet. To accomplish this goal, we continue to make improvements to our shopping cart platform, and we have run periodic incentives to encourage distributors to place their orders via the internet. As a result of these initiatives, approximately 40% of our order volume in the U.S. is placed via internet.

These systems are designed to provide financial and operating data for management, timely and accurate product ordering, generation royalty payment calculation and processing, inventory management, and detailed distributor records. We intend to continue to invest in our systems in order to help meet our business strategies.

Intellectual Property

Our formulas are protected as trade secrets and, to the extent necessary, by confidentiality agreements. In addition, we have obtained U.S. patents on five products as set forth below:

<u>Product</u>	<u>Patent Expiration Date</u>
Innergize!	November 2012
FibRestore	June 2014
Cellebrate	June 2015
Arthaffect	March 2018
ReversAge	May 2021

Currently, we have 22 trademarks registered with the U.S. Patent and Trademark Office, or USPTO, including Reliv and the names of 15 of our 17 nutritional products. NOW for Kids and 24K are not registered with the USPTO. Trademark registrations for selected marks have been issued or applied for in Australia, New Zealand, Canada, Mexico, the United Kingdom, Ireland, the Philippines, Malaysia, Singapore, Germany and several other foreign countries that offer network marketing opportunities. We consider our trademarks to be an important asset of our business.

Regulation

Product Regulation

The formulation, manufacturing, labeling and advertising or promotion of our products are subject to regulation by the Food and Drug Administration, or FDA, which regulates our products under the federal Food, Drug and Cosmetic Act, or FDCA, the Federal Trade Commission, or FTC, and various agencies of the states or countries into which our products are shipped or sold. FDA regulations include requirements and limitations with respect to the labeling of our food and cosmetic products and also with respect to the formulation of those products. FDA regulations also limit and control the extent to which health or other claims can be made with respect to the efficacy of any food or cosmetic. The FDCA has been amended several times with respect to dietary supplements, most recently by the Nutrition Labeling and Education Act of 1990, or NLEA, and the Dietary Supplement Health and Education Act of 1994, or DSHEA, and related regulations. Such legislation governs the formulation, manufacturing, marketing and sale of nutritional supplements, including the content and presentation of health-related information included on the labels or labeling of nutritional supplements.

The majority of the products we market are classified as dietary supplements under the FDCA. Dietary supplements such as those we manufacture and sell, for which no "drug" claim is made, are not subject to FDA approval prior to their sale. However, DSHEA established a pre-market notification process for dietary supplements that contain a "new dietary ingredient," or NDI, a term that is defined as "a dietary ingredient that was not marketed in the United States before October 15, 1994," the date on which DSHEA was signed into law. Certain NDIs that have been "present in the food supply" are exempt from the notification requirement. For those NDIs that are not exempt, DSHEA requires the manufacturer or distributor of a dietary supplement containing an NDI to submit to the FDA, at least 75 days prior to marketing, a notification containing the basis for concluding that the dietary supplement containing the NDI will "reasonably be expected to be safe." Dietary supplement products can be removed from the market if shown to be unsafe, or if the FDA determines, based on the labeling of products, that the intended use of the product is for the diagnosis, cure, mitigation, treatment or prevention of disease. The FDA can regulate those products as "drugs" and require premarket approval of a "new drug application." Manufacturers of

dietary supplements that make any claims for dietary supplements, including product performance and health benefit claims, must have substantiation that the statements are truthful and not misleading.

In January 2000, the FDA published a final rule that defines the types of statements that can be made concerning the effect of a dietary supplement on the structure or function of the body pursuant to the DSHEA. Under the DSHEA, dietary supplement labeling may bear "structure/function" claims, which are claims that the products affect the structure or function of the body, without prior FDA approval. They may not, without prior FDA approval, bear a claim that they can prevent, treat, cure, mitigate or diagnose disease, otherwise known as a "drug claim." The final rule describes how the FDA will distinguish drug claims from structure/function claims. Dietary supplements, like conventional foods, are also permitted to make "health claims," which are claims that are exempt from regulation as "drug" claims pursuant to the amendments to the FDCA established by the NLEA in 1990. A "health claim" is a claim, ordinarily approved by FDA regulation, on a food or dietary supplement product's labeling that "characterizes the relationship of any substance to a disease or health-related condition." To help assure that foods, dietary supplements and cosmetics comply with the provisions of the FDCA and FDA's regulations, the FDA has numerous enforcement tools, including the ability to issue warning letters, initiate product seizures and injunctions and pursue criminal penalties.

The manufacture of dietary supplements is subject to existing FDA current good manufacturing practice, or cGMP, regulations for food. In June 2007, the FDA issued new regulations relating to more detailed cGMP specifically for dietary supplements. Under the new regulations, we qualify as a small business and became subject to the regulations in June 2009. In September 2009 and in February 2011 our Chesterfield plant was audited by the FDA. We received no notice of deviations from cGMP on Form 483 as a result of the 2009 audit and expect that we will receive no notice of deviations from the 2011 audit. We believe our systems and facilities in Chesterfield are in full compliance with cGMP.

Advertisements for our products are subject to regulation by the FTC. The FTC prohibits unfair methods of competition and unfair or deceptive acts or practices in or affecting commerce and provides that the dissemination of any false advertisement pertaining to drugs, cosmetics or foods, including dietary supplements, is an unfair or deceptive practice. Under the FTC's substantiation doctrine, an advertiser must have a "reasonable basis" for all claims made about a product. The failure to be able to adequately substantiate claims may be considered either deceptive or unfair practices. In order to avoid a violation of the FTC standards, we endeavor to assure that we have adequate substantiation for all advertising claims made for our products. In addition, the FTC has increased its scrutiny of the use of distributor testimonials. Although it is impossible for us to monitor all the product claims made by our independent distributors, we make efforts to monitor distributor testimonials and restrict inappropriate distributor claims. The FTC has been more aggressive in pursuing enforcement against dietary supplement products since the passage of DSHEA in 1994, and has brought numerous actions against dietary supplement companies, some resulting in several million dollar civil penalties and/or restitution as well as court-ordered injunctions.

We are aware that, in some of our international markets, there has been recent adverse publicity concerning products that contain substances generally referred to as "genetically modified organisms," or GMOs. In some markets, the possibility of health risks thought to be associated with GMOs has prompted proposed or actual governmental regulation. When necessary, we have responded to government regulations that forbid products containing GMOs by changing certain unacceptable ingredients to non-GMO substitutes. Some of our products in certain markets still contain substances that would be or might be classified as GMOs. We cannot anticipate the extent to which future regulations in these markets will restrict the use of GMOs in our products or the impact of any regulations on our business in those markets. In response to any applicable future regulations, we intend to reformulate our products to satisfy the regulations. Compliance with regulatory requirements in this area should not have a material adverse effect on our business.

Sales Program Regulation

Our distribution and sales program is subject to regulation by the FTC and other federal and state regulation as well as regulations in several countries in which we conduct business. Various state agencies regulate multi-level distribution services. We are required to register with, and submit information to, certain of such agencies and we believe we have complied fully with such requirements. We actively strive to comply with all applicable state and federal laws and regulations affecting our products and our sales and distribution programs. The Attorneys General of several states have taken an active role in investigating and prosecuting companies whose compensation plans

they claim violate local anti-pyramid and/or consumer protection statutes. We are unable to predict the effect such increased activity will have on our business in the future nor are we able to predict the probability of future laws, regulations or interpretations which may be passed by state or federal regulatory authorities.

Federal and state laws directed at network marketing programs have been adopted throughout the years to prevent the use of fraudulent practices often characterized as “pyramid schemes.” Illegal pyramid schemes compensate participants primarily for the introduction or enrollment of additional participants into the program. Often these schemes are characterized by large up-front entry or sign-up fees, over-priced products of low value, little or no emphasis on the sale or use of products, high-pressure recruiting tactics and claims of huge and quick financial rewards with little or no effort. Generally, these laws are directed at ensuring that product sales ultimately are made to consumers and that advancement within such sales organizations is based on sales of products. We have obtained approval of our marketing program as required in all of the markets where we operate and do so for each country we enter.

We believe that our network marketing system satisfies the standards and case law defining a legal marketing system. It is an ongoing part of our business to monitor and respond to regulatory and legal developments, including those that may affect our network marketing system. However, the regulatory and legal requirements concerning network marketing systems do not include “bright line” rules and are inherently fact-based.

Competition

The business of developing and distributing nutritional and skin care products such as those we offer is highly competitive. Numerous manufacturers, distributors and retailers compete for consumers and, in the case of other network marketing companies, for distributors. Our competitors include both network marketing companies such as Alticor Inc. (Amway Corp.), Avon Products Inc., Herbalife Ltd., Mary Kay Inc., Melaleuca, Inc., Mannatech, Inc., Nature’s Sunshine Products Inc., NuSkin Enterprises Inc. and USANA Health Sciences Inc., as well as specialty and mass retail establishments. Our ability to remain competitive depends on the underlying science and high quality of our products and our success in recruiting and retaining distributors. The pool of individuals interested in network marketing tends to be limited in each market and may be reduced to the extent other network marketing companies successfully recruit these individuals into their businesses. We believe that we offer a rewarding compensation plan with attractive financial benefits to compete for the time, attention and commitment of distributors. Our compensation plan is seamless, permitting international expansion.

Reliv NOW and Reliv Classic compete with numerous supplements that offer multi-vitamin benefits. The Reliv Ultrim-Plus, Slimplicity and Celebrate products compete with other products in the weight loss market, including nationally advertised products such as SlimFast. Many companies have entered, or have plans to enter, the sports drink market in which Innergize! and ProVantage compete, a market led by Gatorade. With Arthraffect, FibRestore, ReversAge, GlucAffect, CardioSentials, SoySentials and the Relivables skin care and food products, we are in the specific wellness needs, food and anti-aging markets, which are extremely competitive and led by the major food and skin care companies.

Employees

As of December 31, 2010, we and all of our subsidiaries had approximately 246 full-time employees compared with 239 such employees at the end of 2009.

Additional Available Information

We make available, free of charge, copies of our annual reports on Form 10-K, quarterly reports on Form 10-Q, current reports on Form 8-K, and all amendments to these reports as soon as reasonably practicable after such material is electronically filed with, or furnished to the SEC pursuant to Section 13(a) or 15(d) of the Exchange Act. This information is available on our corporate web site at www.reliv.com under the “Investor Relations” section. This information may also be obtained from the SEC’s on-line database located at www.sec.gov.

Item No. 2 – Properties

We own approximately six acres of land and a building containing approximately 126,000 square feet of office, manufacturing and warehouse space located in Chesterfield, Missouri, where we maintain our corporate headquarters and sole manufacturing facility. We believe that our worldwide facilities are suitable and adequate in relation to our present and immediate future needs.

The following table summarizes information related to our worldwide facilities as of December 31, 2010:

<u>Location</u>	<u>Nature of Use</u>	<u>Square Feet</u>	<u>Owned/Leased</u>
Chesterfield, MO, USA	corporate headquarters/call center/manufacturing/warehouse	126,000	Owned
Seven Hills (Sydney), Australia	central office/warehouse/distribution	8,900	Leased
Oakville, Ontario, Canada	warehouse/distribution	2,100	Leased
Mexico City, Mexico	central office/warehouse/distribution	28,000	Leased
Makati City (Manila), Philippines	central office/warehouse/distribution	3,900	Leased
Birmingham, England, UK	central office/warehouse/distribution	2,200	Leased
Petaling Jaya, Malaysia	central office/call center warehouse/distribution	4,000	Leased
Jakarta, Indonesia	central office/warehouse/distribution	1,600	Leased

Item No. 3 - Legal Proceedings

From time to time, we are involved in litigation incidental to the conduct of our business. We do not believe that any current proceedings will have a material adverse effect on our business, financial condition, results of operations or cash flows.

Item No. 4 - (Removed and Reserved)

PART II

Item No. 5 - Market for Registrant's Common Equity, Related Stockholder Matters and Issuer Purchases of Equity Securities

Our common stock is listed on the NASDAQ Global Select Market under the symbol: RELV. The following table sets forth the high and low sales prices of our common stock and the quarterly dividends per share paid on our common stock during the years ended December 31, 2010 and 2009.

	High	Low	Dividend
Year Ending December 31, 2010			
Fourth Quarter	\$ 2.18	\$ 1.64	\$ 0.02
Third Quarter	2.43	2.00	-
Second Quarter	3.10	2.25	0.02
First Quarter	3.48	2.87	-
Year Ending December 31, 2009			
Fourth Quarter	\$ 3.75	\$ 2.97	\$ 0.02
Third Quarter	4.12	2.85	-
Second Quarter	4.80	2.08	0.05
First Quarter	5.04	3.40	-

As of March 1, 2011, there were approximately 1,685 holders of record of our common stock and an additional 3,653 beneficial owners, including shares of common stock held in street name.

Item No. 7 - Management's Discussion and Analysis of Financial Condition and Results of Operations

The following discussion and analysis of our financial condition and results of operations should be read in conjunction with our financial statements and related notes included elsewhere in this Annual Report on Form 10-K. The following discussion and analysis discusses the financial condition and results of our operations on a consolidated basis, unless otherwise indicated.

Overview

We are a developer, manufacturer and marketer of a proprietary line of nutritional supplements addressing basic nutrition, specific wellness needs, weight management and sports nutrition. We also offer a line of skin care and food products under our Relivables brand. We sell our products through an international network marketing system utilizing independent distributors. Sales in the United States represented approximately 85.0% of worldwide net sales for the year ended December 31, 2010 compared to approximately 87.9% for the year ended December 31, 2009. Our international operations currently generate sales through distributor networks with facilities in Australia, Canada, Indonesia, Malaysia, Mexico, the Philippines, and the United Kingdom. We also operate on a limited basis in Ireland, Germany, Austria and the Netherlands from our U.K. distribution center, in New Zealand from our Australia office, and in Singapore and Brunei from our Malaysia office.

We derive our revenues principally through product sales made by our global independent distributor base, which, as of December 31, 2010, consisted of approximately 60,740 distributors. Our sales can be affected by several factors, including our ability to attract new distributors and retain our existing distributor base, our ability to properly train and motivate our distributor base and our ability to develop new products and successfully maintain our current product line.

All of our sales to distributors outside the United States are made in the respective local currency; therefore, our earnings and cash flows are subject to fluctuations due to changes in foreign currency rates as compared to the U.S. dollar. As a result, exchange rate fluctuations may have an effect on sales and gross margins. Accounting practices require that our results from operations be converted to U.S. dollars for reporting purposes. Consequently, our reported earnings may be significantly affected by fluctuations in currency exchange rates, generally increasing with a weaker U.S. dollar and decreasing with a strengthening U.S. dollar. Products manufactured by us for sale to our foreign subsidiaries are transacted in U.S. dollars. From time to time, we enter into foreign exchange forward contracts to mitigate our foreign currency exchange risk.

Components of Net Sales and Expense

Product sales represent the actual product purchase price typically paid by our distributors, after giving effect to distributor allowances, which can range between 20% to 40% of suggested retail price, depending on the rank of a particular distributor. Handling and freight income represents the amounts billed to distributors for shipping costs. We record net sales and the related commission expense when the merchandise is shipped.

Our primary expenses include cost of products sold, distributor royalties and commissions and selling, general and administrative expenses.

Cost of products sold primarily consists of expenses related to raw materials, labor, quality control and overhead directly associated with production of our products and sales materials, as well as shipping costs relating to the shipment of products to distributors, and duties and taxes associated with product exports. Cost of products sold is impacted by the cost of the ingredients used in our products, the cost of shipping distributors' orders, along with our efficiency in managing the production of our products.

Distributor royalties and commissions are monthly payments made to distributors, based on products sold in their downline organization. Based on our distributor agreements, these expenses typically approximate 23% of sales at suggested retail. Also, we include other sales leadership bonuses, such as Ambassador bonuses, in this line

item. Distributor royalties and commissions are directly related to the level of our sales and, absent any changes in our distributor compensation plan, should continue at comparable levels as a percentage of net sales as in recent periods.

Selling, general and administrative expenses include the compensation and benefits paid to our employees except for those in manufacturing, all other selling expenses, marketing, promotional expenses, travel and other corporate administrative expenses. These other corporate administrative expenses include professional fees, non-manufacturing depreciation and amortization, occupancy costs, communication costs and other similar operating expenses. Selling, general and administrative expenses can be affected by a number of factors, including staffing levels and the cost of providing competitive salaries and benefits; the amount we decide to invest in distributor training and motivational initiatives; and the cost of regulatory compliance.

Results of Operations

The following table sets forth selected results of our operations expressed as a percentage of net sales for the years ended December 31, 2010 and 2009. Our results of operations for the periods described below are not necessarily indicative of results of operations for future periods.

	<u>2010</u>	<u>2009</u>
Net sales	100.0%	100.0%
Costs and expenses:		
Cost of products sold	20.0	19.7
Distributor royalties and commissions..	37.4	37.7
Selling, general and administrative.....	<u>38.9</u>	<u>38.1</u>
Income from operations.....	3.7	4.5
Interest income	0.1	0.1
Interest expense	(0.3)	(0.3)
Other income	<u>0.1</u>	<u>0.4</u>
Income before income taxes	3.6	4.7
Provision for income taxes	<u>1.5</u>	<u>1.7</u>
Net income.....	<u><u>2.1%</u></u>	<u><u>3.0%</u></u>

Year Ended December 31, 2010 Compared to Year Ended December 31, 2009

Net Sales. Overall, sales decreased by 7.8% worldwide, as sales in the United States decreased by 10.9% in the year ended December 31, 2010 compared to 2009. During 2010, our international sales increased by 14.4% over the prior year. All of our international markets, except Asia, showed an increase in sales during 2010 compared to the prior year.

The following table summarizes net sales by geographic market for the years ended December 31, 2010 and 2009.

	Year Ended December 31,					
	2010		2009		Change from prior year	
	Amount	% of Net Sales	Amount	% of Net Sales	Amount	%
	(dollars in thousands)					
United States.....	\$ 66,896	85.0%	\$ 75,041	87.9%	\$ (8,145)	(10.9)%
Australia/New Zealand.....	2,548	3.2	2,459	2.9	89	3.6
Canada.....	2,159	2.8	1,548	1.8	611	39.5
Mexico.....	1,435	1.8	1,371	1.6	64	4.7
Europe.....	2,080	2.6	1,335	1.5	745	55.8
Asia.....	3,630	4.6	3,645	4.3	(15)	(0.4)
Consolidated total.....	\$ 78,748	100.0%	\$ 85,399	100.0%	\$ (6,651)	(7.8)%

The following table sets forth, as of December 31, 2010 and 2009, the number of our active distributors and Master Affiliates and above. The total number of active distributors includes Master Affiliates and above. We define an active distributor as one that enrolls as a distributor or renews its distributorship during the prior twelve months. Master Affiliates and above are distributors that have attained the highest level of discount and are eligible for royalties generated by Master Affiliates and above in their downline organization. Growth in the number of active distributors and Master Affiliates and above is a key factor in continuing the growth of our business.

	December 31, 2010		December 31, 2009		% Change	
	Active Distributors	Master Affiliates and Above	Active Distributors	Master Affiliates and Above	Active Distributors	Master Affiliates and Above
	United States.....	47,450	6,990	54,040	8,640	(12.2)%
Australia/New Zealand.....	2,210	190	2,540	210	(13.0)	(9.5)
Canada.....	1,380	200	1,170	140	17.9	42.9
Mexico.....	1,850	300	2,200	260	(15.9)	15.4
Europe.....	2,170	300	1,150	180	88.7	66.7
Asia.....	5,680	620	6,840	720	(17.0)	(13.9)
Consolidated total.....	60,740	8,600	67,940	10,150	(10.6)%	(15.3)%

Sales in the United States continue to be adversely impacted by distributor uncertainty in the economic recovery and the reduced availability of consumer credit. In addition to the direct impact on sales, these factors lead to fewer distributors qualifying for the level of Master Affiliate. In 2010, approximately 2,040 distributors qualified as new Master Affiliates and 57.3% of the Master Affiliates and above as of December 31, 2009 requalified as Master Affiliates and above during 2010. This compares to approximately 2,570 new Master Affiliates and a requalification rate of 55.6% in 2009. The net number of Master Affiliates and above as of December 31, 2010 decreased by 19.1%, compared to the number as of December 31, 2009.

Another impact to our business due to the downturn in the economy is the average order size. In the United States during 2010, we processed approximately 260,000 orders for products at an average order of \$336 at suggested retail. In 2009, we processed approximately 275,000 product orders at an average order of \$353 at suggested retail. This decline in the average order size is another indicator of the impact of the current economic conditions and a contributing factor in the lower numbers of distributors reaching the Master Affiliate level.

The net number of active Distributors in the United States as of December 31, 2010 decreased by 12.2% to 47,450, compared to the number of active Distributors as of December 31, 2009. Another factor in the decline in sales in the United States was fewer new distributor enrollments. During 2010, approximately 13,620 new distributors were enrolled, compared to 19,580 new distributor enrollments in 2009, a decline of 30.4%. In response to the decline in new distributor enrollments, we indefinitely reduced the cost to enroll as a distributor to \$25 in August 2010. We also revised the materials that a new distributor receives by targeting the materials to entry level distributors. In 2009, we ran an initiative from January through August 2009 to increase new distributor enrollments by offering an enrollment fee of \$20, half of the then-normal \$39.95 fee. Distributor retention in the United States declined slightly to approximately 62.6% for 2010 compared to a rate of 63.1% for 2009. As part of the changes to the distributor enrollment process in August 2010, we also reduced the cost of the annual distributorship renewal from \$30 to \$25.

During the year ended December 31, 2010, net sales in our international operations increased in aggregate by 14.4% to \$11.85 million compared to \$10.36 million for the year ended December 31, 2009. Net sales increased in all of our international markets, except in Asia. Although international sales increased in aggregate during 2010, approximately 58% of the increase was the result of foreign currency fluctuation due to a weaker U.S. dollar. When net sales for the full year of 2010 are converted using the 2009 exchange rate for both 2010 and 2009, international net sales increased by 6.0% for 2010 compared to the prior year. The average exchange rate for the U.S. dollar for all of 2010 was weaker against all currencies of the countries we conduct business, except for the British pound and the euro, compared to the average exchange rates for all of 2009.

Net sales in the Australia/New Zealand market increased by 3.6% in 2010 compared to 2009. New distributor enrollments were 793 in 2010 compared to 952 in 2009. In 2010, 49 distributors qualified as new Master Affiliates, compared to 67 in the prior year. When net sales are converted using the 2009 exchange rate for both 2010 and 2009, net sales in this market decreased by 11.6%. Net income for the Australia/New Zealand market was \$33,000 in 2010, compared to net income of \$7,000 in 2009.

Net sales in Canada increased by 39.5% in 2010 compared to 2009. When measured in local currency, Canadian net sales increased by 25.9% in 2010 compared to 2009, as we had a strong increase in the number of new Master Affiliate qualifications. In 2010, 88 distributors qualified as new Master Affiliates, compared to 48 in the prior year. New distributor enrollments were 548 in 2010 compared to 476 in 2009. Net loss in Canada was \$15,000 for 2010, compared to a net income of \$35,000 in 2009. The improvement in gross profit was offset by a reduction in foreign currency transaction gains. For all of 2010, we recorded gains of \$39,000, compared to transaction gains of \$126,000 for 2009.

Net sales in Mexico increased 4.7% in 2010 compared to 2009. New distributor enrollments were 1,306 in 2010 compared to 1,700 in 2009, and 194 distributors qualified as new Master Affiliates in 2010, compared to 140 in the prior year. When measured in local currency, 2010 net sales decreased by 2.1%, as the Mexican peso strengthened on average for 2010 when compared to the U.S. dollar. The net loss in Mexico for 2010 was \$300,000, compared to a net loss of \$316,000 in 2009.

Our European region includes sales from operations in United Kingdom, Ireland, Germany, Austria and the Netherlands. Net sales in Europe increased by 55.8% for 2010 compared to 2009. When measured in local currency, net sales in Europe increased by 58.8% in 2010 compared to the prior year. Strong company sales leadership, coupled with improving local distributor leadership led to the increase. New distributor enrollments were 1,518 in 2010 compared to 617 in 2009, and 188 distributors qualified as new Master Affiliates in 2010, compared to 96 in 2009. The net loss incurred in Europe was \$320,000 in 2010, compared to a net loss of \$478,000 in 2009. The 2010 net loss was reduced as a result of the increase in current year sales.

Our Asian region includes sales from operations from the Philippines, Malaysia, Singapore, Brunei, and Indonesia. Net sales in Asia decreased by 0.4% in 2010 compared to the prior year. New distributor enrollments were 3,864 in 2010 compared to 4,941 in 2009, and 326 distributors qualified as new Master Affiliates in 2010, compared to 336 in 2009. When measured in local currency, 2010 net sales decreased by 6.8%. The net loss in Asia for 2010 was \$664,000, compared to a net loss of \$437,000 in 2009, as the result of decline in sales. Asian sales declined, particularly in the second and third quarters of 2010, as the result of departures in our local management staff in our Malaysian office and declining trends in both new distributor enrollments and new Master

Affiliate qualifications. We hired a new general manager late in the second quarter of 2010 to oversee the Southeast Asia area, which includes our Malaysia, Singapore, Brunei, and Indonesia operations.

Cost of Products Sold. Cost of products sold as a percentage of net sales increased to 20.0% for the year ended December 31, 2010 compared to 19.7% for the year ended December 31, 2009. Gross margins declined in 2010 compared to 2009 due to changes in revenue mix and lower production levels corresponding with the decrease in sales.

Distributor Royalties and Commissions. Distributor royalties and commissions as a percentage of net sales decreased slightly to 37.4% for the year ended December 31, 2010 compared to 37.7% for the same period in 2009. The minor reduction is due to the introduction of the Relivables product line in the third quarter of 2009, which has a lower commission rate.

Selling, General and Administrative Expenses. For 2010, selling, general and administrative, or SGA, expenses decreased by \$1.91 million compared to 2009. However, SGA expenses as a percentage of net sales increased to 38.9% in 2010 compared to 38.1% in 2009, as a function of the 7.8% decline in consolidated net sales.

Sales and marketing expenses decreased by \$1.15 million in 2010. Of that amount, \$395,000 represented the decrease in expenses directly related to sales volume, such as star director bonuses, other sales production bonuses, and credit card fees. Other changes included a decrease of \$391,000 for our distributor conferences and other training events, and a decrease of \$123,000 for distributor newsletter costs.

General and administrative expenses, including salaries and benefits, decreased by approximately \$727,000 in 2010 compared to 2009. One of the significant decreases was in salaries, incentive compensation expense and benefits of \$562,000, as we reduced the contribution to our Employee Stock Ownership Plan by \$340,000 for the year. Other significant changes included a decrease in legal, accounting and consulting fees of \$153,000; business insurance expenses of \$100,000; and depreciation expense of \$141,000. Amortization expense increased by \$137,000, related to intangible assets associated with the acquisition of a Reliv distributorship in 2009.

Interest Income/Expense. Interest income decreased to \$48,000 for the year ended December 31, 2010, compared to \$52,000 for the same period in 2009. The decrease in interest income is the result of lower interest rates during 2010. Interest expense increased to \$206,000 for 2010 compared to \$174,000 for 2009, as the result of a full year of interest expense for two long-term debt agreements we entered into during 2009.

Income Taxes. We recorded income tax expense of \$1.14 million for 2010, representing an effective rate of 40.4%. In 2009, we recorded income tax expense of \$1.47 million, representing an effective rate of 36.9%. The higher effective rate in 2010 is the result of the increased impact of the permanent non-deductible expenses compared to pre-tax income and a higher effective rate of state income taxes.

Net Income. Our net income decreased to \$1.68 million (\$0.14 per share basic and diluted) for the year ended December 31, 2010 compared to \$2.52 million (\$0.20 per share basic and diluted) for 2009. Profitability decreased commensurate with the decrease in net sales in the United States, as discussed above, plus a slight increase in the net loss from international operations. Net income in the United States was \$2.95 million in 2010, compared to \$3.70 million in 2009. The net loss from international operations was \$1.27 million in 2010, compared a net loss of \$1.19 million in 2009.

Financial Condition, Liquidity and Capital Resources

We generated \$2.23 million of net cash during 2010 from operating activities, \$845,000 was used in investing activities, and we used \$1.02 million in financing activities. This compares to \$5.76 million of net cash provided by operating activities, \$1.01 million used in investing activities, and \$3.65 million used in financing activities in 2009. Cash and cash equivalents increased by \$570,000 to \$6.33 million as of December 31, 2010 compared to December 31, 2009.

Significant changes in working capital items consisted of an increase in inventory of \$472,000, and a decrease in accounts payable and accrued expenses of \$535,000 in 2010. The increase in inventory is the result of lower than expected sales levels, compared to scheduled production. We also increased inventory levels in some of

our foreign markets to provide better lead time and shipping efficiencies. The decrease in accounts payable and accrued expenses is related to a lower level of payables with production vendors. Furthermore, accrued distributor commission expense is approximately \$263,000 lower at the end of 2010 compared to the end of 2009.

Our net investing activities included \$577,000 and \$534,000 in net capital expenditures for the years ended December 31, 2010 and 2009, respectively. Payments for key-man life insurance were \$268,000 in 2010 and \$252,000 in 2009. Other investing activities in 2009 also included \$716,000 in cash payment for the purchase of a distributorship, along with proceeds of \$489,000 from the final withdrawal in a limited partnership investment.

Financing activities in 2010 consisted of \$521,000 in payments on long-term debt and \$495,000 in common stock dividends paid. Financing activities in 2009 included \$5.01 million in payments for purchases of our common stock into treasury and \$856,000 in common stock dividends paid.

Stockholders' equity increased to \$13.93 million at December 31, 2010 compared with \$12.27 million at December 31, 2009. The increase represents our net income of \$1.68 million for 2010, offset by our cash dividend of \$495,000. Other changes to equity include the contribution of treasury shares to our ESOP of \$125,000, an improvement in our cumulative foreign currency translation adjustment of \$180,000, and other equity-based compensation of \$171,000.

Our working capital balance was \$6.86 million at December 31, 2010 compared to \$5.47 million at December 31, 2009. The current ratio at December 31, 2010 was 2.07 compared to 1.81 at previous year-end.

On November 30, 2010, we entered into a term loan with our primary lender ("the Bank") in the principal amount of \$3.66 million. The loan was renegotiated from a loan that originated with the Bank on June 29, 2009. The term of the loan is for a period of three years with interest accruing on the outstanding principal balance at a floating interest rate based on the 30-day LIBOR plus 2.0%. Monthly principal and interest payments are based on approximately a nine-year amortization. The aggregate outstanding balance of principal and interest is due and payable on November 30, 2013.

We also renewed a revolving credit facility for \$5 million with the Bank. The credit facility accrues interest on the outstanding principal balance at a floating interest rate based on 30-day LIBOR plus 1.85% and has a maturity date of September 30, 2011. As of December 31, 2010, there were no outstanding borrowings on the revolving credit facility.

The amended terms of the term loan and revolving credit facility are reflected in separate promissory notes dated November 30, 2010 between us and the Bank. A separate letter agreement dated June 29, 2009 stating the financial covenants related to the term loan and revolving credit facility continues in effect.

Under the terms of the letter agreement, we have agreed to financial covenants under which we are required to (i) maintain at all times a tangible net worth of not less than \$10 million and (ii) maintain at all times a ratio of Total Funded Debt to EBITDA of not greater than 2.5 to 1. The term loan and revolving credit facility are secured by all of our tangible and intangible assets and also by a mortgage on our building and real estate located in Chesterfield, Missouri. As of December 31, 2010, we were in compliance with all financial covenants.

Management believes that our internally generated funds coupled with the bank loan facilities will be sufficient to meet working capital requirements for the remainder of 2011.

Critical Accounting Policies

Our financial statements are based on the selection and application of significant accounting policies, which require management to make significant estimates and assumptions. We believe that the following are some of the more critical judgment areas in the application of our accounting policies that currently affect our financial condition and results of operations.

Revenue

We receive payment by credit card, personal check, or guaranteed funds for orders from independent distributors and make related commission payments in the following month. Net sales reflect product sales at suggested retail price less the distributor discount of 20% to 40%. Sales revenue and commission expenses are recorded when the merchandise is shipped, as this is the point title and risk of loss pass. In accordance with FASB ASC, Topic 650-50, "Revenue Recognition-Customer Payments and Incentives," we present distributor royalty and commission expense as an operating expense, rather than a reduction to net sales, as these payments are not made to the purchasing distributor.

Actual and estimated returns are classified as a reduction of net sales. We estimate and accrue a reserve for product returns based on our return policy and historical experience. Our return policy allows for distributors to return product only upon termination of his or her distributorship. Allowable returns are limited to saleable product which was purchased within twelve months of the termination for a refund of 90% of the original purchase price less any distributor royalties and commission received relating to the original purchase of the returned products. Total returns have been approximately 0.67% and 0.70% of net sales in 2010 and 2009, respectively. We record handling and freight income as a component of net sales and record handling and freight costs as a component of cost of products sold. Total revenues do not include sales tax as we consider ourselves a pass-through conduit for collecting and remitting applicable sales taxes.

Inventories

Inventories are valued at the lower of cost or market. Product cost includes raw material, labor and overhead costs and is accounted for using the first-in, first-out basis. On a periodic basis, we review our inventory levels in each country for estimated obsolescence or unmarketable items, as compared to future demand requirements and the shelf life of the various products. Based on this review, we record inventory write-downs when costs exceed expected net realizable value. Historically, our estimates of obsolete or unmarketable items have been materially accurate.

Sales aids and promotional materials inventories represent distributor kits, product brochures, and other sales and business development materials which are held for sale to distributors. Costs of the sales aids and promotional materials held for sale are capitalized as inventories and subsequently recorded to cost of goods sold upon recognition of revenue when sold to distributors. All other advertising and promotional costs are expensed when incurred.

Foreign Currency Translation

All balance sheet accounts are translated using the exchange rates in effect at the balance sheet date. Statements of operations amounts are translated using the average exchange rate for the year-to-date periods. The gains and losses resulting from the changes in exchange rates during the period have been reported in other comprehensive loss. Foreign currency translation adjustments exclude income tax expense (benefit) given that our investments in non-U.S. subsidiaries are deemed to be reinvested for an indefinite period of time.

Legal Proceedings

In the ordinary course of business, we are subject to various legal proceedings, including lawsuits and other claims related to labor, product and other matters. We are required to assess the likelihood of adverse judgments and outcomes to these matters as well as the range of potential loss. Such assessments are required to determine whether a loss contingency reserve is required under the provisions of FASB ASC Topic 450, "Contingencies," and to determine the amount of required reserves, if any. These assessments are subjective in nature. Management makes these assessments for each individual matter based on consultation with outside counsel and based on prior experience with similar claims. To the extent additional information becomes available or our strategies or assessments change, our estimates of potential liability for a given matter may change. Changes to estimates of liability would result in a corresponding additional charge or benefit recognized in the statement of operations in the period in which such changes become known. We recognize the costs associated with legal defense in the periods incurred. Accordingly, the future costs of defending claims are not included in our estimated liability.

Stock-Based Compensation

We have stock-based incentive plans under which we may grant stock option, restricted stock, and unrestricted stock awards. We recognize stock-based compensation expense based on the grant date fair value of the award and the related vesting terms as proscribed in FASB ASC Topic 718, "Compensation-Stock Compensation." We use the Black-Scholes option pricing model to determine the fair value of stock options which requires us to estimate certain key assumptions. For the years ended December 31, 2010 and 2009, we incurred employee stock-based compensation cost of \$190,000 (\$124,000 net of tax), and \$192,000 (\$126,000 net of tax), respectively.

Income Tax Matters

We account for income taxes in accordance with FASB ASC Topic 740, "Income Taxes," (ASC Topic 740) which requires that deferred tax assets and liabilities be recognized using enacted tax rates for the effect of temporary differences between the book and tax bases of recorded assets and liabilities. ASC Topic 740 also requires that deferred tax assets be reduced by a valuation allowance if it is "more likely than not" that some portion or the entire deferred tax asset will not be realized. In our quarterly evaluation of the need for a valuation allowance, we take into account various factors, including the expected level of future taxable income and available tax planning strategies. If actual results differ from the assumptions made in our previous evaluation of our valuation allowance, we may record a change in valuation allowance through income tax expense in the period this determination is made.

At December 31, 2010, we had deferred tax assets related to net operating loss carryforwards and other income tax credits with a tax value of \$4.3 million. These net operating loss carryforwards have various expiration dates, depending on the country and period in which they occurred. A valuation allowance of \$4.3 million has been established for these deferred tax assets based on projected future taxable income and the expiration dates of these carryforwards.

At December 31, 2010, we also had deferred tax assets related to 2008 capital losses on investments with a tax value of \$349,000. We have established a corresponding valuation allowance of \$349,000 against this deferred tax asset as we do not anticipate having sufficient future capital gains to offset these capital losses.

The calculations of our tax liabilities involve dealing with uncertainties in the application of complex tax regulations. On January 1, 2007, we adopted provisions of ASC Topic 740 related to uncertain tax positions. As a result of the implementation of the provisions, we recognize liabilities for uncertain tax positions based on the two-step process prescribed in the guidance. The first step is to evaluate the tax position for recognition by determining if the weight of available evidence indicates that it is more likely than not that the position will be sustained on audit, including resolution of related appeals or litigation processes, if any. The second step requires us to estimate and measure the tax benefit as the largest amount that is more than 50% likely to be realized upon ultimate settlement. It is inherently difficult and subjective to estimate such amounts, as we have to determine the probability of various possible outcomes. We reevaluate these uncertain tax positions on a quarterly basis. This evaluation is based on factors including, but not limited to, changes in facts or circumstances, changes in tax law, effectively settled issues under audit, or new audit activity. Such a change in recognition or measurement would result in the recognition of a tax benefit or an additional charge to the tax provision.

Item No. 8 - Financial Statements and Supplementary Data

Reference is made to the Consolidated Financial Statements contained in Part IV hereof.

Item No. 9 - Changes in and Disagreements with Accountants on Accounting and Financial Disclosure

None

Item No. 9A - Controls and Procedures

Evaluation of Disclosure Controls and Procedures

Our management, under the supervision and with the participation of our Chief Executive Officer and Chief Financial Officer, has reviewed and evaluated the effectiveness of the design and operation of our disclosure controls and procedures as of December 31, 2010. Based on such review and evaluation, our Chief Executive Officer and Chief Financial Officer have concluded that the disclosure controls and procedures were effective as of December 31, 2010, to ensure that the information required to be disclosed by us in the reports that we file or submit under the Securities Exchange Act of 1934, as amended, (a) is recorded, processed, summarized and reported within the time period specified in the SEC's rules and forms and (b) is accumulated and communicated to our management, including the officers, as appropriate to allow timely decisions regarding required disclosure.

Management's Annual Report on Internal Control over Financial Reporting

Management is responsible for establishing and maintaining adequate internal control over financial reporting. Our management conducted an evaluation of the effectiveness of our internal control over financial reporting based on the framework in Internal Control – Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission. This evaluation included review of the documentation of controls, evaluation of the design effectiveness of controls, testing of the operation effectiveness of controls and a conclusion on this evaluation. Although there are inherent limitations in the effectiveness of any system of internal control over financial reporting, based on our evaluation, management has concluded our internal controls over financial reporting were effective as of December 31, 2010.

Attestation Report of the Registered Public Accounting Firm

This annual report does not include an attestation report of the company's registered public accounting firm regarding internal control over financial reporting. Management's report was not subject to attestation by the company's registered public accounting firm as the company is classified as a "Smaller Reporting Company."

Changes in Internal Control over Financial Reporting

There were no material changes in our internal control over financial reporting during the fourth quarter of 2010 that have materially affected or are reasonably likely to materially affect our internal controls over financial reporting.

Item No. 9B - Other Information

None

PART III

Item No. 10 - Directors and Executive Officers of the Registrant

Information called for by Item 10 of Part III is incorporated by reference to the definitive Proxy Statement for the 2011 Annual Meeting of Shareholders to be held on May 26, 2011, which is expected to be filed with the Commission within 120 days after December 31, 2010.

Item No. 11 - Executive Compensation

Information called for by Item 11 of Part III is incorporated by reference to the definitive Proxy Statement for the 2011 Annual Meeting of Shareholders to be held on May 26, 2011, which is expected to be filed with the Commission within 120 days after December 31, 2010.

Item No. 12 - Security Ownership of Certain Beneficial Owners and Management and Related Stockholder Matters

Information called for by Item 12 of Part III is incorporated by reference to the definitive Proxy Statement for the 2011 Annual Meeting of Shareholders to be held on May 26, 2011, which is expected to be filed with the Commission within 120 days after December 31, 2010.

Item No. 13 - Certain Relationships and Related Transactions

Information called for by Item 13 of Part III is incorporated by reference to the definitive Proxy Statement for the 2011 Annual Meeting of Shareholders to be held on May 26, 2011, which is expected to be filed with the Commission within 120 days after December 31, 2010.

Item No. 14 - Principal Accountant Fees and Services

Information called for by Item 14 of Part III is incorporated by reference to the definitive Proxy Statement for the 2011 Annual Meeting of Shareholders to be held on May 26, 2011, which is expected to be filed with the Commission within 120 days after December 31, 2010.

PART IV

Item No. 15 - Exhibits and Financial Statement Schedules

- (a) 1. The Consolidated Financial Statements filed as part of this report on Form 10-K are listed on the accompanying Index to Consolidated Financial Statements and Consolidated Financial Statement Schedules.
2. Financial schedules required to be filed by Item 8 of this form, and by Item 15(d) below:
- All other financial schedules are not required under the related instructions or are inapplicable and therefore have been omitted.
3. Exhibits:

**Exhibit
Number**

Document

- 3.1 Second Amended and Restated Certificate of Incorporation (incorporated by reference to Appendix B of Schedule 14A of the Registrant filed on April 17, 2003).
- 3.2 By-Laws (incorporated by reference to the Registration Statement on Form S-3 of the Registrant filed on February 21, 2006).
- 3.3 Amendment to By-Laws dated March 22, 2001 (incorporated by reference to the Registration Statement on Form S-3 of the Registrant filed on February 21, 2006).
- 3.4 Certificate of Designation to Create a Class of Series A Preferred Stock for Reliv' International, Inc. (incorporated by reference to Exhibit 3.1 to the Form 10-Q of the Registrant for quarter ended March 31, 2003).
- 4.1 Form of Reliv International, Inc. common stock certificate (incorporated by reference to the Registration Statement on Form S-3 of the Registrant filed on February 21, 2006).
- 10.1 Amended Exclusive License Agreement with Theodore P. Kalogris dated December 1, 1991 (incorporated by reference to Exhibit 10.1 to the Form 10-K of the Registrant for the year ended December 31, 1992).

- 10.2* Robert L. Montgomery Employment Agreement dated June 19, 2007 (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed June 25, 2007).
- 10.3* Carl W. Hastings Employment Agreement dated July 26, 2007 (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed July 27, 2007).
- 10.4 Letter Agreement with Southwest Bank of St. Louis dated June 29, 2009 (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed July 6, 2009).
- 10.5 Promissory Note (Term Loan) dated November 30, 2010 by the Registrant in favor of M&I Marshall and Ilsley Bank (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed December 6, 2010).
- 10.6 Promissory Note (Revolving Credit Facility) dated November 30, 2010 by the Registrant in favor of M&I Marshall and Ilsley Bank (incorporated by reference to Exhibit 10.2 to the Form 8-K of the Registrant filed December 6, 2010).
- 10.7* Reliv' International, Inc. Supplemental Executive Retirement Plan dated June 1, 1998 (incorporated by reference to Exhibit 10.19 to the Form 10-K of the Registrant for year ended December 31, 1998).
- 10.8* Reliv International, Inc. Employee Stock Ownership Plan and Trust dated August 24, 2006 (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed August 30, 2006).
- 10.9* 2009 Distributor Stock Purchase Plan (incorporated by reference to Appendix 1 of Form S-3 Registration Statement the Registrant filed July 1, 2009).
- 10.10* 2003 Stock Option Plan (incorporated by reference to Exhibit 4 to the Form S-8 Registration Statement the Registrant filed August 13, 2003).
- 10.11* 2009 Incentive Stock Plan (incorporated by reference to Exhibit 10.1 to the Form S-8 Registration Statement the Registrant filed December 2, 2010).
- 10.12* Reliv International, Inc. Incentive Compensation Plan effective January 1, 2007 (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed May 31, 2007).
- 10.13* R. Scott Montgomery Employment Agreement dated January 2, 2008 (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed January 4, 2008).
- 10.14* Ryan A. Montgomery Employment Agreement dated January 2, 2008 (incorporated by reference to Exhibit 10.2 to the Form 8-K of the Registrant filed January 4, 2008).
- 10.15* Steven G. Hastings Employment Agreement dated January 2, 2008 (incorporated by reference to Exhibit 10.3 to the Form 8-K of the Registrant filed January 4, 2008).
- 10.16* Steven D. Albright Employment Agreement dated January 2, 2008 (incorporated by reference to Exhibit 10.4 to the Form 8-K of the Registrant filed January 4, 2008).
- 10.17* Brett M. Hastings Employment Agreement dated January 2, 2008 (incorporated by reference to Exhibit 10.5 to the Form 8-K of the Registrant filed January 4, 2008).
- 10.18 Stock Purchase Agreement among the Paul and Jane Meyer Family Foundation and Reliv International, Inc. dated April 23, 2009 (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed April 28, 2009).

- 10.19 Purchase Agreement by and among Michael G. Williams, Julie T. Williams and Reliv International, Inc. dated August 31, 2009 (incorporated by reference to Exhibit 10.1 to the Form 8-K of the Registrant filed September 3, 2009).
- 11 Statement re: computation of per share earnings (incorporated by reference to Note 8 of the Consolidated Financial Statements contained in Part IV).
- 21 Subsidiaries of the Registrant (filed herewith).
- 23 Consent of Ernst & Young LLP, Independent Auditors (filed herewith).
- 31.1 Certification of Chief Executive Officer pursuant to Rule 13a-14(a) and Rule 15d-14(a) of the Securities Exchange Act, as amended (filed herewith).
- 31.2 Certification of Chief Financial Officer pursuant to Rule 13a-14(a) and Rule 15d-14(a) of the Securities Exchange Act, as amended (filed herewith).
- 32 Certification of Chief Executive Officer and Chief Financial Officer pursuant to 18 U.S.C. 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002 (filed herewith).

*Indicates management compensation plan, contract or arrangement.

By: /s/ Michael D. Smith
Michael D. Smith, Director

Date: March 17, 2011

By: /s/ John M. Klimek
John M. Klimek, Director

Date: March 17, 2011

Exhibit Index

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Reliv' International, Inc.
and Subsidiaries

Consolidated Financial Statements

Years ended December 31, 2010 and 2009

Contents

Consolidated Financial Statements:

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Consolidated Statements of Stockholders' Equity for the years ended December 31, 2010 and 2009	F-5
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Report of Independent Registered Public Accounting Firm

The Board of Directors and Stockholders
Reliv' International, Inc.

We have audited the accompanying consolidated balance sheets of Reliv' International, Inc. and Subsidiaries (the Company) as of December 31, 2010 and 2009, and the related consolidated statements of income, stockholders' equity, and cash flows for each of the two years in the period ended December 31, 2010. These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements based on our audits.

We conducted our audits in accordance with the standards of the Public Company Accounting Oversight Board (United States). Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. We were not engaged to perform an audit of the Company's internal control over financial reporting. Our audits included consideration of internal control over financial reporting as a basis for designing audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the Company's internal control over financial reporting. Accordingly, we express no such opinion. An audit includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements. An audit also includes assessing the accounting principles used and significant estimates made by management, as well as evaluating the overall financial statement presentation. We believe that our audits provide a reasonable basis for our opinion.

In our opinion, the financial statements referred to above present fairly, in all material respects, the consolidated financial position of Reliv' International, Inc. and Subsidiaries at December 31, 2010 and 2009, and the consolidated results of their operations and their cash flows for each of the two years in the period ended December 31, 2010, in conformity with U.S. generally accepted accounting principles.

/s/ Ernst & Young LLP

St. Louis, Missouri
March 17, 2011

Reliv' International, Inc. and Subsidiaries

Consolidated Balance Sheets

	December 31	
	2010	2009
Assets		
Current assets:		
Cash and cash equivalents	\$ 6,331,038	\$ 5,760,913
Accounts receivable, less allowances of \$67,100 in 2010 and \$59,700 in 2009	291,405	326,022
Accounts due from employees and distributors	55,854	78,500
Inventories:		
Finished goods	3,851,178	3,073,570
Raw materials	1,277,838	1,388,140
Sales aids and promotional materials	521,774	622,694
Total inventories	5,650,790	5,084,404
Refundable income taxes	62,324	23,789
Prepaid expenses and other current assets	519,915	652,544
Deferred income taxes	334,000	303,000
Total current assets	13,245,326	12,229,172
Other assets	364,626	333,279
Cash surrender value of life insurance	1,503,350	1,235,800
Intangible assets, net	1,785,987	1,991,497
Property, plant, and equipment	18,980,656	18,629,377
Less accumulated depreciation	11,036,244	10,264,692
	<u>7,944,412</u>	<u>8,364,685</u>
Total assets	<u>\$ 24,843,701</u>	<u>\$ 24,154,433</u>

Reliv' International, Inc. and Subsidiaries

Consolidated Balance Sheets (continued)

	December 31	
	2010	2009
Liabilities and stockholders' equity		
Current liabilities:		
Accounts payable and accrued expenses	\$ 5,820,291	\$ 6,242,289
Current maturities of long-term debt	566,873	519,192
Total current liabilities	<u>6,387,164</u>	<u>6,761,481</u>
Noncurrent liabilities:		
Long-term debt, less current maturities	4,150,770	4,719,542
Other noncurrent liabilities	375,244	406,544
Total noncurrent liabilities	<u>4,526,014</u>	<u>5,126,086</u>
Stockholders' equity:		
Preferred stock, par value \$0.001 per share; 3,000,000 shares authorized; -0- shares issued and outstanding in 2010 and 2009	-	-
Common stock, par value \$0.001 per share; 30,000,000 shares authorized, 14,425,185 shares issued and 12,450,808 shares outstanding in 2010; 14,425,185 shares issued and 12,380,187 shares outstanding in 2009	14,425	14,425
Additional paid-in capital	30,300,463	30,228,573
Accumulated deficit	(10,091,167)	(11,279,526)
Accumulated other comprehensive loss:		
Foreign currency translation adjustment	(448,024)	(627,704)
Treasury stock	(5,845,174)	(6,068,902)
Total stockholders' equity	<u>13,930,523</u>	<u>12,266,866</u>
Total liabilities and stockholders' equity	<u>\$ 24,843,701</u>	<u>\$ 24,154,433</u>

See accompanying notes.

Reliv' International, Inc. and Subsidiaries

Consolidated Statements of Income

	Year ended December 31	
	2010	2009
Product sales	\$ 70,033,580	\$ 75,845,599
Handling & freight income	8,714,808	9,553,471
Net sales	<u>78,748,388</u>	<u>85,399,070</u>
Costs and expenses:		
Cost of products sold	15,738,885	16,862,622
Distributor royalties and commissions	29,450,171	32,172,148
Selling, general, and administrative	<u>30,652,045</u>	<u>32,557,704</u>
Income from operations	<u>2,907,287</u>	<u>3,806,596</u>
Other income (expense):		
Interest income	47,744	52,292
Interest expense	(205,985)	(173,867)
Other income	<u>76,110</u>	<u>300,260</u>
Income before income taxes	<u>2,825,156</u>	<u>3,985,281</u>
Provision for income taxes	<u>1,142,000</u>	<u>1,470,000</u>
Net income available to common shareholders	<u>\$ 1,683,156</u>	<u>\$ 2,515,281</u>
Earnings per common share - Basic	<u>\$0.14</u>	<u>\$0.20</u>
Weighted average shares	<u>12,382,000</u>	<u>12,894,000</u>
Earnings per common share - Diluted	<u>\$0.14</u>	<u>\$0.20</u>
Weighted average shares	<u>12,383,000</u>	<u>12,894,000</u>

See accompanying notes.

Reliv' International, Inc. and Subsidiaries
Consolidated Statements of Stockholders' Equity

	Common Stock		Additional Paid-In Capital	Accumulated Deficit	Accumulated Other Comprehensive Loss	Treasury Stock		Total
	Shares	Amount				Shares	Amount	
Balance at December 31, 2008	14,425,185	\$ 14,425	\$ 30,321,066	\$ (12,938,430)	\$ (663,478)	123,025	\$ (625,993)	\$ 16,107,590
Net income	-	-	-	2,515,281	-	-	-	2,515,281
Other comprehensive income:								
Foreign currency translation adjustment	-	-	-	-	35,774	-	-	35,774
Total comprehensive income	-	-	-	-	-	-	-	2,551,055
Common stock dividends paid, \$0.07 per share	-	-	-	(856,377)	-	-	-	(856,377)
Stock-based compensation, net of excess tax benefits	-	-	120,632	-	-	-	-	120,632
Contribution of treasury shares to ESOP	-	-	(213,125)	-	-	(150,000)	678,125	465,000
Common stock purchased for treasury	-	-	-	-	-	2,071,973	(6,121,034)	(6,121,034)
Balance at December 31, 2009	14,425,185	14,425	30,228,573	(11,279,526)	(627,704)	2,044,998	(6,068,902)	12,266,866
Net income	-	-	-	1,683,156	-	-	-	1,683,156
Other comprehensive income:								
Foreign currency translation adjustment	-	-	-	-	179,680	-	-	179,680
Total comprehensive income	-	-	-	-	-	-	-	1,862,836
Common stock dividends paid, \$0.04 per share	-	-	-	(495,344)	-	-	-	(495,344)
Stock-based compensation, net of excess tax benefits	-	-	170,618	-	-	-	-	170,618
Contribution of treasury shares to ESOP	-	-	(98,728)	-	-	(70,621)	223,728	125,000
Other	-	-	-	547	-	-	-	547
Balance at December 31, 2010	14,425,185	\$ 14,425	\$ 30,300,463	\$ (10,091,167)	\$ (448,024)	1,974,377	\$ (5,845,174)	\$ 13,930,523

See accompanying notes.

Reliv' International, Inc. and Subsidiaries

Consolidated Statements of Cash Flows

	Year ended December 31	
	2010	2009
Operating activities		
Net income	\$ 1,683,156	\$ 2,515,281
Adjustments to reconcile net income to net cash provided by operating activities:		
Depreciation and amortization	1,244,990	1,192,948
Stock-based compensation	170,618	120,632
Contribution of treasury shares to ESOP	125,000	465,000
Deferred income taxes	(67,000)	52,000
Foreign currency transaction (gain)/loss	(69,592)	(147,606)
(Increase) decrease in accounts receivable	72,862	505,788
(Increase) decrease in inventories	(471,532)	1,237,953
(Increase) decrease in refundable income taxes	(38,833)	107,269
(Increase) decrease in prepaid expenses and other current assets	145,133	349,214
(Increase) decrease in other assets	(31,818)	(35,827)
Increase (decrease) in accounts payable & accrued expenses and other non-current liabilities	(535,328)	(603,771)
Net cash provided by operating activities	<u>2,227,656</u>	<u>5,758,881</u>
Investing activities		
Proceeds from sale of property, plant, and equipment	41,332	3,978
Purchase of property, plant, and equipment	(618,545)	(537,617)
Payment of life insurance premiums	(267,550)	(252,200)
Purchase of distributorship	-	(716,119)
Proceeds from final withdrawal from limited partnership investment	-	488,633
Net cash used in investing activities	<u>(844,763)</u>	<u>(1,013,325)</u>
Financing activities		
Proceeds from line of credit borrowings	-	6,000,000
Repayment of line of credit borrowings	-	(6,000,000)
Proceeds from term loan borrowings	-	4,120,000
Principal payments on short and long-term borrowings	(521,091)	(1,901,442)
Common stock dividends paid	(495,344)	(856,377)
Purchase of stock for treasury	-	(5,014,115)
Other	547	-
Net cash used in financing activities	<u>(1,015,888)</u>	<u>(3,651,934)</u>
Effect of exchange rate changes on cash and cash equivalents	203,120	206,654
Increase (decrease) in cash and cash equivalents	<u>570,125</u>	<u>1,300,276</u>
Cash and cash equivalents at beginning of year	<u>5,760,913</u>	<u>4,460,637</u>
Cash and cash equivalents at end of year	<u>\$ 6,331,038</u>	<u>\$ 5,760,913</u>

Reliv' International, Inc. and Subsidiaries

Consolidated Statements of Cash Flows (continued)

	Year ended December 31	
	2010	2009
Supplemental disclosures of cash flow information:		
Cash paid during the year for:		
Interest	\$ 206,356	\$ 177,200
Income taxes	\$ 1,304,000	\$ 1,378,000
Noncash investing and financing transactions:		
Issuance of promissory notes for purchase of stock for treasury	\$ -	\$ 1,106,919
Obligation for purchase of distributorship	\$ -	\$ 1,343,881

See accompanying notes.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

December 31, 2010

1. Nature of Business and Significant Accounting Policies

Nature of Business

Reliv' International, Inc. (the Company) produces a proprietary line of nutritional supplements addressing basic nutrition, specific wellness needs, weight management, and sports nutrition. These products are sold by subsidiaries of the Company to a sales force of independent distributors and licensees of the Company that sell products directly to consumers. The Company and its subsidiaries sell products to distributors throughout the United States and in Australia, Austria, Brunei, Canada, Germany, Indonesia, Ireland, Malaysia, Mexico, the Netherlands, New Zealand, the Philippines, Singapore, and the United Kingdom.

Principles of Consolidation

The consolidated financial statements include the accounts of the Company and its foreign and domestic subsidiaries. All significant intercompany accounts and transactions have been eliminated.

Inventories

Inventories are valued at the lower of cost or market. Product cost includes raw materials, labor, and overhead costs and is accounted for on a first-in, first-out basis. On a periodic basis, the Company reviews its inventory levels, as compared to future demand requirements and the shelf life of the various products. Based on this review, the Company records inventory write-downs when necessary.

Sales aids and promotional materials inventories represent distributor kits, product brochures, and other sales and business development materials which are held for sale to distributors. Cost of the sales aids and promotional materials held for sale are capitalized as inventories and subsequently recorded to cost of goods sold upon recognition of revenue when sold to distributors. All other advertising and promotional costs are expensed when incurred.

Property, Plant, and Equipment

Property, plant, and equipment are stated on the cost basis. Depreciation is computed using the straight-line or an accelerated method over the useful life of the related assets. Generally, computer equipment and software are depreciated over 5 years, office equipment and machinery over 7 years, and real property over 39 years.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

1. Nature of Business and Significant Accounting Policies (continued)

Foreign Currency Translation and Transaction Gains or Losses

All balance sheet accounts have been translated using the exchange rates in effect at the balance sheet date. Statements of income amounts have been translated using the average exchange rate for the year. The gains and losses resulting from the changes in exchange rates from year to year have been reported in other comprehensive income (loss). The foreign currency translation adjustment is the only component of accumulated other comprehensive loss. If applicable, foreign currency translation adjustments exclude income tax expense (benefit) as certain of the Company's investments in non-U.S. subsidiaries are deemed to be reinvested for an indefinite period of time. Transaction gains were \$69,592 and \$147,606 for 2010 and 2009, respectively.

Revenue Recognition

The Company receives payment by credit card, personal check, or guaranteed funds for orders from independent distributors and makes related commission payments in the following month. Generally, net sales reflect product sales less the distributor discount of 20 percent to 40 percent of the suggested retail price. Sales revenue and commission expenses are recorded when the merchandise is shipped, as this is the point title and risk of loss pass to the distributor. In accordance with Financial Accounting Standards Board ("FASB") Accounting Standards Codification ("ASC") Topic 605-50, "Revenue Recognition – Customer Payments and Incentives," the Company presents distributor royalty and commission expense as an operating expense, rather than a reduction to net sales, as these payments are not made to the purchasing distributor.

Actual and estimated sales returns are classified as a reduction of net sales. The Company estimates and accrues a reserve for product returns based on the Company's return policy and historical experience. The Company's return policy allows for distributors to return product only upon termination of his or her distributorship. Allowable returns are limited to saleable product which was purchased within twelve months of the termination for a refund of 90% of the original purchase price less any distributor royalties and commission received relating to the original purchase of the returned products. For the years ended December 31, 2010 and 2009, total returns as a percent of net sales were approximately 0.67 % and 0.70%, respectively.

The Company records handling and freight income as a component of net sales and records handling and freight costs as a component of cost of products sold. Total revenues do not include sales tax as the Company considers itself a pass-through conduit for collecting and remitting applicable sales taxes.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

1. Nature of Business and Significant Accounting Policies (continued)

Basic and Diluted Earnings per Share

Basic earnings per common share are computed using the weighted average number of common shares outstanding during the year. Diluted earnings per common share are computed using the weighted average number of common shares and potential dilutive common shares that were outstanding during the period. Potential dilutive common shares consist of outstanding stock options, outstanding stock warrants, and convertible preferred stock. See Note 8 for additional information regarding earnings per share.

Stock-Based Compensation

The Company has stock-based incentive plans under which it may grant stock option, restricted stock, and unrestricted stock awards. The Company recognizes stock-based compensation expense based on the grant date fair value of the award and the related vesting terms. The fair value of stock-based awards is determined using the Black-Scholes model, which incorporates assumptions regarding the risk-free interest rate, expected volatility, expected option life, and dividend yield. See Note 7 for additional information.

The Company accounts for options granted to non-employees and warrants granted to distributors under the fair value approach required by FASB ASC Topic 505-50, "Equity Based Payments to Non-Employees."

Income Taxes

The provision for income taxes is computed using the liability method. The primary differences between financial statement and taxable income result from financial statement accruals and reserves and differences between depreciation for book and tax purposes.

Unrecognized tax benefits are accounted for as required by FASB ASC Topic 740 which prescribes a more likely than not threshold for financial statement presentation and measurement of a tax position taken or expected to be taken in a tax return. See Note 11 for further discussion.

Fair Value Measurements

FASB ASC Topic 820, "Fair Value Measurements and Disclosures," defines fair value, establishes a framework for measuring fair value, and requires disclosures about fair value measurements required under other accounting pronouncements. See Note 5 for further discussion.

Advertising

Costs of sales aids and promotional materials are capitalized as inventories. All other advertising and promotional costs are expensed when incurred. The Company recorded \$55,000 and \$91,000 of advertising expense in 2010 and 2009, respectively.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

1. Nature of Business and Significant Accounting Policies (continued)

Amortizable Intangible Assets

The Company records intangible assets based on management's determination of the fair value of the respective assets at the time of acquisition. Determining the fair value of intangible assets is judgmental and involves the use of significant estimates and assumptions of future company operations. The Company bases its fair value estimates and related asset lives on assumptions it believes to be reasonable but that are unpredictable and inherently uncertain. Actual future results may differ from these estimates.

Intangible assets estimated to have finite estimable lives are amortized over their estimated economic life under the straight-line method. Based on management's estimates, these lives range from two to fifteen years. Related amortization expense is presented within Selling, General, and Administrative in the accompanying consolidated statements of income. See Note 14 for further information.

Research and Development Expenses

Research and development expenses, which are charged to selling, general, and administrative expenses as incurred, were \$587,000 and \$551,000 in 2010 and 2009, respectively.

Cash Equivalents

The Company's policy is to consider the following as cash and cash equivalents: demand deposits, short-term investments with a maturity of three months or less when purchased, and highly liquid debt securities with both insignificant interest rate risk and with original maturities from the date of purchase of generally three months or less.

Use of Estimates

The preparation of financial statements in conformity with accounting principles generally accepted in the United States requires management to make estimates and assumptions that affect the amounts reported in the financial statements and accompanying notes. Actual results could differ from those estimates.

Reclassifications

At December 31, 2010, the cash surrender value of life insurance balance exceeds 5% of total assets and is therefore separately presented in the consolidated balance sheets and statements of cash flows. To conform to the 2010 presentation, 2009 amounts have been reclassified from Other Assets.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

2. Property, Plant, and Equipment

Property, plant, and equipment at December 31, 2010 and 2009, consist of the following:

	<u>2010</u>	<u>2009</u>
Land and land improvements	\$ 868,870	\$ 852,147
Building	9,928,950	9,851,829
Machinery and equipment	3,698,537	3,426,720
Office equipment	1,503,929	1,494,915
Computer equipment and software	2,980,370	3,003,766
	<u>18,980,656</u>	<u>18,629,377</u>
Less accumulated depreciation	11,036,244	10,264,692
	<u>\$ 7,944,412</u>	<u>\$ 8,364,685</u>

3. Accounts Payable and Accrued Expenses

Accounts payable and accrued expenses at December 31, 2010 and 2009, consist of the following:

	<u>2010</u>	<u>2009</u>
Trade payables	\$ 2,437,965	\$ 2,627,674
Distributors' commissions	2,411,016	2,674,247
Sales taxes	445,653	362,612
Payroll and payroll taxes	525,657	577,756
	<u>\$ 5,820,291</u>	<u>\$ 6,242,289</u>

4. Limited Partnership Investment

In June 2006, the Company contributed \$1,000,000 as a limited partner in a private equity fund. In accordance with FASB ASC Topic 323-30, "Investments – Equity Methods and Joint Ventures," the Company accounted for its investment under the equity method. Under this method, the Company's proportionate share of partnership income or loss was recorded to gain or loss on limited partnership investment with a corresponding increase or decrease in the carrying value of its investment.

In 2008, the Company delivered notice to the partnership's general partner of its notice to fully withdraw from the partnership. Therefore, the carrying value of the investment at December 31, 2008 of \$489,000 was included in "Prepaid Expenses and Other Current Assets" in the consolidated balance sheets. Upon final cessation of the partnership in 2009, the Company received cash from the partnership of \$489,000.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

5. Fair Value of Financial Instruments

The fair value of financial instruments at December 31, 2010 and 2009 were as follows:

Description	Fair Value	Level 1	Level 2	Level 3
<u>December 31, 2010</u>				
Long-term debt	\$4,613,000	-	\$4,613,000	-
Marketable securities ⁽¹⁾	232,000	\$232,000	-	-
Derivatives ⁽²⁾	11,693	-	11,693	-
<u>December 31, 2009</u>				
Long-term debt	\$5,184,000	-	\$5,184,000	-
Marketable securities ⁽¹⁾	200,000	200,000	-	-

(1) Representing assets of the Company's Supplemental Executive Retirement Plan (trading securities). Presented within Other Assets in the consolidated balance sheets.

(2) Representing recorded liability of Canadian forward currency contracts and is presented within Accounts Payable and Accrued Expenses in the consolidated balance sheets. The fair values of derivatives are determined either through quoted market prices in active markets for exchange traded derivatives or through pricing from brokers who develop values based on inputs observable in active markets such as interest rates and currency volatilities.

Fair value can be measured using valuation techniques such as the market approach (comparable market prices), the income approach (present value of future income or cash flow), and the cost approach (cost to replace the service capacity of an asset or replacement cost). Accounting standards utilize a fair value hierarchy that prioritizes the inputs to valuation techniques used to measure fair value into three broad levels. The following is a brief description of those three levels:

Level 1: Observable inputs such as quoted prices (unadjusted) in active markets for identical assets or liabilities.

Level 2: Inputs other than quoted prices that are observable for the asset or liability, either directly or indirectly. These include quoted prices for similar assets or liabilities in active markets or similar assets or liabilities in markets that are not active.

Level 3: Unobservable inputs that reflect the reporting entity's own assumptions.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

5. Fair Value of Financial Instruments (continued)

At December 31, the carrying amount and fair value of the Company's financial instruments are approximately as follows:

	2010		2009	
	Carrying Amount	Fair Value	Carrying Amount	Fair Value
Long-term debt	\$4,717,643	\$4,613,000	\$5,238,734	\$5,184,000
Marketable securities	232,000	232,000	200,000	200,000
Derivatives	11,693	11,693	-	-

The carrying value of other financial instruments, including cash, accounts receivable and accounts payable, and accrued liabilities approximate fair value due to their short maturities or variable-rate nature of the respective balances.

6. Long-Term Debt

Long-term debt at December 31, 2010 and 2009 consists of the following:

	2010	2009
Term loan	\$ 3,594,908	\$ 3,948,768
Obligation for purchase of distributorship	1,122,735	1,289,966
	<u>4,717,643</u>	<u>5,238,734</u>
Less current maturities	566,873	519,192
	<u>\$ 4,150,770</u>	<u>\$ 4,719,542</u>

Principal maturities of long-term debt at December 31, 2010, are as follows:

2011	\$ 566,873
2012	584,741
2013	2,998,096
2014	204,171
2015	214,617
Thereafter	149,145
	<u>\$ 4,717,643</u>

Effective October 1, 2008, the Company entered into a \$5 million revolving loan agreement (2008) for a one year term with its primary lender. Effective October 1, 2009, upon expiration of the 2008 revolving loan agreement, the Company entered into a new \$5 million one-year revolving loan agreement (2009) with its primary lender.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

6. Long-Term Debt (continued)

Effective November 30, 2010, the Company entered into a new one-year \$5 million revolving loan agreement (2010) with its primary lender. Any advances under the revolver accrue interest at a variable interest rate based on 30-day LIBOR + 1.85%. Interest, if any, is payable monthly. At December 31, 2010, the outstanding revolving line of credit balance was zero.

2009 Purchases of Stock for Treasury and related Borrowings

In April 2009, the Company entered into a Stock Purchase Agreement with a significant shareholder (Seller) to purchase 2,068,973 shares of the Company's common stock for \$6,106,919 (an average price of \$2.95 per share). To finance the purchase, the Company borrowed \$5 million under its 2008 revolving line of credit and issued a promissory note to the Seller for \$1,106,919. The promissory note bore interest at 6% per annum with all principal and interest due no later than ninety days from closing. The Company repaid this promissory note in July 2009 by borrowing \$1 million from its 2008 revolving line of credit.

In June 2009, the Company entered into a term loan agreement with its primary lender for \$4.12 million and used all of the proceeds to reduce its revolving line of credit balance. The term loan was for a period of two years with interest accruing at a floating interest rate based on the 30-day LIBOR plus 3%, subject to a 3.75% floor. Monthly principal and interest were based on a ten-year amortization. As originally structured, the aggregate outstanding balance of principal and interest was due and payable on June 29, 2011.

On November 30, 2010, the Company re-financed its term loan agreement with its primary lender. The re-financed term loan is for a period of three years with interest accruing at a floating interest rate based on the 30-day LIBOR plus 2%. As of December 31, 2010, the term loan's interest rate was 2.27%. Monthly principal and interest are based on approximately a nine-year amortization. The aggregate outstanding balance of principal and interest is due and payable on November 30, 2013.

The term loan agreement and 2010 revolving line of credit agreement are secured by all tangible and intangible assets of the Company and also by a mortgage on the real estate of the Company's headquarters. These agreements also include loan covenants requiring the Company to maintain net tangible worth of not less than \$10 million, and that borrowings under the agreements shall not exceed EBITDA by a ratio of 2.5:1. At December 31, 2010, the Company was in compliance with its loan covenants.

Obligation for Purchase of Distributorship

As described in Note 14, on August 31, 2009, the Company incurred a long-term obligation of \$1,343,881 relating to the purchase of a Reliv distributorship. The Company will pay this obligation in monthly payments of principal and interest totaling \$18,994 over a seven-year term with an annual interest rate of 5%.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

7. Stockholders' Equity

Stock Options

2009 Incentive Stock Plan

The Company sponsors an incentive stock plan (the "2009 Plan") allowing for a maximum of 1,000,000 shares to be granted in the form of either incentive stock options, non-qualified stock options, restricted stock awards, or unrestricted stock awards. Employees, directors, advisors, and consultants of the Company are eligible to receive the grants. The plan has been approved by the stockholders of the Company. The Compensation Committee of the Board of Directors administers the plan.

The 2009 Plan provides that options may be issued under the plan at an option price not less than fair market value of the stock at the time the option is granted. Under the 2009 Plan, restricted stock of the Company may be granted at no cost to the grantee. The grantees are entitled to dividends and voting rights for their respective shares. Restrictions limit the sale or transfer of these shares during the requisite service period. In addition, the committee may grant or sell unrestricted stock at a purchase price to be determined by the committee.

Vesting terms and restrictions, if applicable, under the plan, are set by the committee and will be 10 years or less. The 2009 Plan expires in 2019. As of December 31, 2010, there were no grants under the 2009 Plan.

2003 Stock Option Plan

The Company sponsors a stock option plan (the "2003 Plan") allowing for incentive stock options and non-qualified stock options to be granted to employees and eligible directors. The plan has been approved by the stockholders of the Company. The 2003 Plan provided that a maximum of 1,000,000 shares may be issued under the plan at an option price not less than the fair market value of the stock at the time the option is granted. The options vest pursuant to the schedule set forth for the plan. With stockholder approval of the 2009 Incentive Stock Plan, the Board of Directors resolved not to award any additional stock option grants under the 2003 Plan.

In 2005, the Company issued grants of 543,000 shares under the 2003 Plan. The 2005 option grants were issued with an exercise price equal to the fair value of the shares at the time of grant and were fully vested in the year of grant. These option grants precede the Company's 2006 adoption of FASB ASC Topic 718, "Compensation-Stock Compensation." Accordingly, no stock-based compensation expense has been recognized relating to the 2005 option grants.

In August 2007, the Company granted options to purchase 216,000 shares of common stock under the 2003 Plan. The options were issued with an exercise price of \$9.74 which was equal to the fair value of the shares at the time of grant.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

7. Stockholders' Equity (continued)

Stock Options (continued)

The fair value of the options granted in 2007 were estimated at the date of grant using a Black-Scholes option pricing model with the following weighted average assumptions: risk-free interest rate of 5.01%; dividend yield of 1.00%; volatility factor of the expected price of the Company's stock of 0.472; an expected life of 4.5 years and a grant date fair value of \$4.07 per share. The options have a term of five years and vest in increments of 25% beginning August 7, 2009 and ending May 1, 2012. Expense for stock options granted in 2007 is recognized on a straight-line basis separately for each vesting portion of the stock option award.

During 2008, the Company granted options to purchase 16,500 and 25,000 shares of common stock with exercise prices of \$5.28 per share and \$5.50 per share, respectively, and a grant-date fair value of \$1.84 per share and \$1.91 per share, respectively. The options' exercise prices were equal to the fair value of the shares at the time of the grant.

The fair value of the options granted in 2008 were estimated at the date of grant using a Black-Scholes option pricing model with the following weighted average assumptions: risk-free interest rate of approximately 3.0%; dividend yield of 1.9%; volatility factor of the expected price of the Company's stock of 0.447; and an expected life of 4.5 years. The options have a term of five years and vest in various increments ranging from one year to 4.67 years.

Compensation cost for the stock option plans was approximately \$190,000 (\$124,000 net of tax) and \$192,000 (\$126,000 net of tax) for the years ended December 31, 2010 and 2009, respectively, and has been recorded in selling, general, and administrative expense. As of December 31, 2010, the total remaining unrecognized compensation cost related to non-vested stock options totaled \$240,000 (\$156,000 net of tax), which will be amortized over the weighted remaining requisite service period of 1.4 years.

A summary of the Company's stock option activity and related information for the years ended December 31 follows:

	2010		2009	
	Options	Weighted Avg. Exercise Price	Options	Weighted Avg. Exercise Price
Outstanding beginning of the year	754,000	\$8.29	763,000	\$8.31
Granted	-		-	
Exercised	-		-	
Forfeited	(1,000)	9.74	(9,000)	9.74
Outstanding at end of year	753,000	\$8.29	754,000	\$8.29
Exercisable at end of year	642,375	\$8.12	576,875	\$8.06

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

7. Stockholders' Equity (continued)

Stock Options (continued)

Range of Exercise Prices	As of December 31, 2010					
	Options Outstanding			Options Exercisable		
	Number Outstanding	Weighted Avg. Remaining Life	Weighted Avg. Exercise Price	Number Exercisable	Weighted Avg. Remaining Life	Weighted Avg. Exercise Price
\$5.28 - \$5.50	41,500	2.67	\$5.41	29,125	2.67	\$5.47
\$7.92	485,000	4.00	7.92	485,000	4.00	7.92
\$8.68	30,000	4.79	8.68	30,000	4.79	8.68
\$9.74	196,500	1.58	9.74	98,250	1.58	9.74
\$5.28 - \$9.74	<u>753,000</u>	3.33	\$8.29	<u>642,375</u>	3.61	\$8.12

The aggregate intrinsic value of stock options outstanding and currently exercisable at December 31, 2010 was \$-0-. Intrinsic value for stock options is calculated based on the exercise price of the underlying awards as compared to the quoted price of the Company's common stock as of the reporting date.

For the years ended December 31, 2010 and 2009, no stock options were exercised.

Distributor Stock Purchase Plan

In November 1998, the Company established a Distributor Stock Purchase Plan (1998 Plan). The plan allows distributors who have reached the "Ambassador" status the opportunity to allocate up to 10% of their monthly compensation into the plan to be used to purchase the Company's common stock at the current market value. The plan also states that at the end of each year, the Company will grant warrants to purchase additional shares of the Company's common stock based on the number of shares purchased by the distributors under the plan during the year. The warrant exercise price will equal the market price for the Company's common stock at the date of issuance. The warrants issued shall be in the amount of 25% of the total shares purchased under the plan during the year. This plan commenced in January 1999, and a total of 26,134 warrants were issued during the year ended December 31, 2008. The warrants are fully vested upon grant. The weighted average fair value of warrants granted during 2008 was \$1.30 per share.

The 1998 Plan was established with a ten-year life. As a result, there will be no further grants from this Plan. Upon exercise, forfeiture or expiration of all outstanding warrants, the 1998 Plan will terminate.

In July 2009, the Company established a new Distributor Stock Purchase Plan (2009 Plan) to replace the expired 1998 Plan. The 2009 Plan, which is similar to the 1998 Plan, commenced in August 2009. A total of 3,179 warrants, at a fair value of \$1.06 per share, were issued on December 31, 2009. A total of 13,684 warrants, at a fair value of \$0.65 per share, were issued on December 31, 2010. The warrants are fully vested upon grant.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

7. Stockholders' Equity (continued)

Distributor Stock Purchase Plan (continued)

The Company records expense under the fair value method for warrants granted to distributors. Total expense recorded for these warrants was \$3,370 and \$-0- in 2010 and 2009, respectively.

The fair value of the warrants was estimated at the date of grant using a Black-Scholes option pricing model with the following assumptions:

	Year ended December 31	
	2010	2009
Expected warrant life (years)	3.0	2.5
Risk-free weighted average interest rate	1.02%	1.70%
Stock price volatility	0.549	0.540
Dividend yield	2.1%	1.2%

A summary of the Company's warrant activity and related information for the years ended December 31 follows:

	2010		2009	
	Warrants	Weighted Avg. Exercise Price	Warrants	Weighted Avg. Exercise Price
Outstanding beginning of the year	53,689	\$6.25	79,040	\$7.25
Granted	13,684	1.94	3,179	3.28
Exercised	-	-	-	-
Expired and forfeited	(25,891)	8.19	(28,530)	8.68
Outstanding at end of year	<u>41,482</u>	\$3.62	<u>53,689</u>	\$6.25
Exercisable at end of year	<u>41,482</u>		<u>53,689</u>	

As of December 31, 2010					
Warrants Outstanding				Warrants Exercisable	
Range of Exercise Prices	Number Outstanding	Weighted Avg. Remaining Life	Weighted Avg. Exercise Price	Number Exercisable	Weighted Avg. Exercise Price
\$ 1.94	13,684	3.00	\$1.94	13,684	\$1.94
\$ 3.28	3,179	2.00	3.28	3,179	3.28
\$ 4.60	24,619	0.88	4.60	24,619	4.60
\$1.94 - \$4.60	<u>41,482</u>	1.67	\$3.62	<u>41,482</u>	\$3.62

The intrinsic value for stock warrants outstanding at December 31, 2010 was \$6,900 with a weighted average remaining life of 1.67 years. For the years ended December 31, 2010 and 2009, no stock warrants were exercised.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

8. Earnings per Share

The following table sets forth the computation of basic and diluted earnings per share:

	Year ended December 31	
	2010	2009
Numerator:		
Net income	\$1,683,156	\$2,515,281
Denominator:		
Denominator for basic earnings per share – weighted average shares	12,382,000	12,894,000
Dilutive effect of employee stock options and other warrants	1,000	-
Denominator for diluted earnings per share – adjusted weighted average shares	12,383,000	12,894,000
Basic earnings per share	\$0.14	\$0.20
Diluted earnings per share	\$0.14	\$0.20

For the year ended December 31, 2010, options and warrants totaling 780,798 shares of common stock were not included in the denominator for diluted earnings per share because their effect would be anti-dilutive. For the year ended December 31, 2009, options and warrants totaling 804,510 shares of common stock were not included in the denominator for diluted earnings per share because their effect would be anti-dilutive.

9. Leases

The Company leases certain office facilities, storage, and equipment. These leases have varying terms, and certain leases have renewal and/or purchase options. Future minimum payments under non-cancelable leases with initial or remaining terms in excess of one year consist of the following at December 31, 2010:

2011	\$ 386,862
2012	232,107
2013	129,197
2014	118,554
2015	61,316
Thereafter	-
	\$ 928,036

Rent expense for operating leases was \$578,281 and \$510,550 for the years ended December 31, 2010 and 2009, respectively.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

10. Derivative Financial Instruments

The Company has various transactions with its foreign subsidiaries that are denominated in U.S. dollars and are thereby subject to foreign currency exchange risk on these transactions.

The Company from time to time uses foreign currency exchange contracts to reduce its exposure to fluctuations in foreign exchange rates. The Company bases these contracts on the amount of cash flows that it expects to be remitted to the United States from its foreign operations and does not use such derivative financial instruments for trading or speculative purposes. The Company accounts for these contracts as free standing derivatives, such that gains or losses on the fair market value of these forward exchange contracts as of the balance sheet dates are recorded as other income and expense in the consolidated statements of income.

At December 31, 2010, the Company held Canadian forward exchange contracts with maturities of less than one year totaling \$487,000. The increase (decrease) in the aggregate accrued loss on these contracts was \$11,692 and \$-0- for the years ended December 31, 2010 and 2009, respectively. No contracts were held at December 31, 2009.

11. Income Taxes

The components of income (loss) before income taxes are as follows:

	Year ended December 31	
	2010	2009
United States	\$4,780,065	\$5,828,091
Foreign	(1,954,909)	(1,842,810)
	\$2,825,156	\$3,985,281

The components of the provision for income taxes are as follows:

	Year ended December 31	
	2010	2009
Current:		
Federal	\$1,041,000	\$1,236,000
State	177,000	185,000
Foreign	33,000	33,000
Total current	1,251,000	1,454,000
Deferred:		
Federal	(93,000)	14,000
State	(16,000)	2,000
Foreign	-	-
Total deferred	(109,000)	16,000
	\$1,142,000	\$1,470,000

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

11. Income Taxes (continued)

The provision for income taxes is different from the amounts computed by applying the United States federal statutory income tax rate of 34%. The reasons for these differences are as follows:

	Year ended December 31	
	2010	2009
Income taxes at U.S. statutory rate	\$961,000	\$1,355,000
State income taxes, net of federal benefit	164,000	204,000
Higher/(lower) effective taxes on earnings in foreign countries	38,000	(13,000)
Foreign corporate income taxes	33,000	33,000
Nondeductible meals and entertainment expense	29,000	35,000
Qualified production activities income - AJCA	(43,000)	(44,000)
Other	(40,000)	(100,000)
	<u>\$1,142,000</u>	<u>\$1,470,000</u>

The components of the deferred tax assets and liabilities, and the related tax effects of each temporary difference at December 31, 2010 and 2009, are as follows:

	2010	2009
Deferred tax assets:		
Product refund reserve	\$ 65,000	\$ 92,000
Inventory obsolescence reserve	26,000	12,000
Vacation accrual	32,000	29,000
Stock-based compensation	237,000	189,000
Organization costs	151,000	91,000
Deferred compensation	80,000	64,000
Capital losses on investments	349,000	343,000
Valuation allowance - investment losses	(349,000)	(343,000)
Miscellaneous accrued expenses	82,000	52,000
Foreign net operating loss carryforwards	4,279,000	3,935,000
Valuation allowance - NOL carryforwards	(4,279,000)	(3,935,000)
	<u>673,000</u>	<u>529,000</u>
Deferred tax liabilities:		
Depreciation	206,000	129,000
Net deferred tax assets (liabilities)	<u>\$ 467,000</u>	<u>\$ 400,000</u>
Reported as:		
Current deferred tax assets	\$ 334,000	\$ 303,000
Non-current deferred tax assets ⁽¹⁾	133,000	97,000
Net deferred tax assets	<u>\$ 467,000</u>	<u>\$ 400,000</u>

(1) Included within other non-current assets on the consolidated balance sheets.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

11. Income Taxes (continued)

The Company has a deferred tax asset of \$4,279,000 as of December 31, 2010, and \$3,935,000 as of December 31, 2009, relating to foreign net operating loss carryforwards. The Company has recorded a valuation allowance as it is more likely than not that this asset will not be realized before it expires beginning in 2011.

The Company has a deferred tax asset as of December 31, 2010 related primarily to 2008 capital losses on investments with a tax value of \$349,000. The Company has established a corresponding valuation allowance of \$349,000 as it does not anticipate having sufficient future capital gains to offset these capital losses. The capital loss carryforward expires in 2013.

Through December 31, 2010, the Company has not recorded a provision for income taxes on the earnings of several of its foreign subsidiaries because such earnings are intended to be permanently reinvested outside the U.S. The cumulative amount of unremitted earnings on which the Company has not recognized United States income tax was \$111,000 at December 31, 2010. Although it is not practicable to determine the deferred tax liability on the unremitted earnings, credits for foreign income taxes paid would be available to significantly reduce any U.S tax liability if foreign earnings are remitted.

The Company's effective tax rate is based on expected income, statutory tax rates and tax planning opportunities available to the Company in the various jurisdictions in which the Company operates. Significant judgment is required in determining the Company's effective tax rate and in evaluating its tax positions. In evaluating the exposure associated with various filing positions, the Company estimates reserves for probable exposures, which are adjusted quarterly in light of changing facts and circumstances, such as the progress of tax audits, case law and emerging legislation.

The Company applied applicable accounting guidance relating to accounting for uncertainty in income taxes. The primary difference between gross unrecognized tax benefits and net unrecognized tax benefits is the U.S. federal tax benefit from state tax deductions. It is the Company's practice to recognize interest and / or penalties related to income tax matters in income tax expense.

At December 31, 2010 and 2009, the Company had \$110,000 and \$166,800, respectively, of cumulative unrecognized tax benefits, all of which would impact the effective income tax rate if recognized.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

11. Income Taxes (continued)

The aggregate changes in the balance of gross unrecognized tax benefits were as follows:

Beginning balance as of January 1, 2009	\$ 160,200
Settlements and effective settlements with tax authorities	(1,400)
Lapse of statute of limitations	-
Increases in balances related to tax positions taken during prior periods	-
Decreases in balances related to tax positions taken during prior periods	(15,000)
Increases in balances related to tax positions taken during current period	23,000
Balance as of December 31, 2009	<u>\$ 166,800</u>
Settlements and effective settlements with tax authorities	(47,800)
Lapse of statute of limitations	-
Increases in balances related to tax positions taken during prior periods	-
Decreases in balances related to tax positions taken during prior periods	(23,000)
Increases in balances related to tax positions taken during current period	14,000
Balance as of December 31, 2010	<u><u>\$ 110,000</u></u>

The current portion of the Company's unrecognized tax benefits is presented in the balance sheet within current liabilities and the amount expected to be settled after one year is recorded in other non-current liabilities.

The Company, including its domestic and foreign subsidiaries, is subject to U.S. federal income tax as well as income tax of multiple state and foreign jurisdictions. The Company has concluded all U.S. federal income tax matters for years through 2006 and concluded years through 2006 with its primary state jurisdiction.

One of the Company's foreign subsidiaries is presently under local country audit for alleged deficiencies in value-added tax (VAT) and withholding tax for the years 2004 through 2006. The Company, in consultation with its legal counsel, believes that there are strong legal grounds that it should not be liable to pay the majority of the alleged tax deficiencies. As of December 31, 2010, management has estimated and reserved approximately \$185,000 for resolution of this matter and recorded this amount within Selling, General, and Administrative expense in the 2010 Consolidated Statements of Income.

12. Employee Benefit Plans

The Company sponsors a 401(k) employee savings plan which covers substantially all employees. Employees can contribute up to 15% of their gross income to the plan, and the Company matches a percentage of the employee's contribution at a rate of 25%. Company contributions under the 401(k) plan totaled \$163,000 and \$147,000 in 2010 and 2009, respectively.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

12. Employee Benefit Plans (continued)

On September 1, 2006, the Company established an employee stock ownership plan ("ESOP") which covers substantially all U.S. employees. Contributions to the ESOP are funded by the Company on a discretionary basis. In 2010 and 2009, the Company's contribution consisted of shares of common stock from treasury measured by the fair value of the stock on date of contribution. Company contributions under the ESOP plan totaled approximately \$125,000 and \$465,000 for the years ended December 31, 2010 and 2009, respectively.

13. Incentive Compensation Plans

In May 2007, the Board of Directors approved the adoption of a new incentive compensation plan. This new plan was effective for fiscal year 2007 and replaced a previous plan. Under the plan, bonuses are payable quarterly in an amount not to exceed 18% of the Company's Income from Operations for any period, subject to the Company achieving a minimum quarterly Income from Operations of at least \$500,000. For fiscal years 2010 and 2009, the Board determined that the aggregate amount of incentive compensation available under the Plan shall be equal to 16% of the Company's Income from Operations. The bonus pool is allocated to executives according to a specified formula, with a portion allocated to a middle management group determined by the Executive Committee of the Board of Directors.

The Company expensed a total of \$457,000 and \$723,000 to the participants of the bonus pool for 2010 and 2009, respectively.

The Company sponsors a Supplemental Executive Retirement Plan (SERP) to allow certain executives to defer a portion of their annual salary and bonus into a grantor trust. A grantor trust was established to hold the assets of the SERP. The Company funds the grantor trust by paying the amount deferred by the participant into the trust at the time of deferral. Investment earnings and losses accrue to the benefit or detriment of the participants. The SERP also provides for a discretionary matching contribution by the Company not to exceed 100% of the participant's annual contribution. In 2010 and 2009, the Company did not provide a match. The participants fully vest in the deferred compensation three years from the date they enter the SERP. The participants are not eligible to receive distribution under the SERP until retirement, death, or disability of the participant. At December 31, 2010 and 2009, SERP assets were \$232,000 and \$200,000, respectively, and are included in "Other Assets" in the accompanying consolidated balance sheets. At December 31, 2010 and 2009, SERP liabilities were \$238,000 and \$211,000, respectively, and are included in "Other Non-Current Liabilities" in the accompanying consolidated balance sheets. The changes in the balances of SERP assets and SERP liabilities during 2010 and 2009 were due to net realized and unrealized investment gains/losses incurred by the plan.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

14. Purchase of Reliv Distributorship

On August 31, 2009, the Company acquired an independent Reliv distributorship from its owner for an aggregate purchase price of \$2,060,000. The Company paid \$500,000 of the purchase price to the owner at closing, credited the owner's \$216,119 outstanding loan balance due to the Company, and will pay the balance of the purchase price, \$1,343,881, over a period of seven years, plus interest at an annual rate of 5%, with monthly payments of principal and interest totaling \$18,994. As a condition to the transaction, the contract contains a non-compete clause of two years and non-solicitation clause of Company distributors for a term of seven years.

The Company allocated the purchase price to its components based on the relative fair values of assets acquired, accounting for the acquisition of the distributorship as an intangible asset with an estimated value of \$1,648,000 and useful life of fifteen years. For the non-compete provision and non-solicitation provision, the Company allocated \$103,000 and \$309,000 respectively, based upon these assets relative fair value estimates and their respective contractual terms.

The distributorship, non-compete, and non-solicit assets, net of accumulated amortization, are presented as "Intangible assets, net" in the accompanying consolidated balance sheets and are subject to review for potential impairment going forward.

The Company had amortizable intangible assets as follows as of December 31, 2010 and 2009:

	<u>Gross Carrying Amount</u>		<u>Accumulated Amortization</u>	
	<u>2010</u>	<u>2009</u>	<u>2010</u>	<u>2009</u>
Distributorship	\$1,648,000	1,648,000	\$146,489	36,622
Non-compete agreement	103,000	103,000	68,667	17,167
Non-solicitation agreement	309,000	309,000	58,857	14,714
	<u>\$2,060,000</u>	<u>2,060,000</u>	<u>\$274,013</u>	<u>68,503</u>

Amortization expense (straight-line method) for intangible assets totaled \$205,510 and \$68,503 in 2010 and 2009, respectively. Amortization expense for amortizable intangible assets over the next five years is estimated to be:

	<u>Intangible Amortization</u>
2011	\$188,300
2012	154,000
2013	154,000
2014	154,000
2015	154,000

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

15. Segment Information

Description of Products and Services by Segment

The Company operates in one reportable segment, a network marketing segment consisting of six operating units that sell nutritional and dietary products to a sales force of independent distributors that sell the products directly to customers. These operating units are based on geographic regions.

Geographic area data for the years ended December 31, 2010 and 2009 follow:

	<u>2010</u>	<u>2009</u>
Net sales to external customers		
United States	\$66,895,693	\$75,041,461
Australia/New Zealand	2,548,046	2,458,834
Canada	2,158,699	1,548,086
Mexico	1,435,486	1,371,127
Europe ⁽¹⁾	2,079,941	1,334,908
Asia ⁽²⁾	3,630,523	3,644,654
Total net sales	<u>\$78,748,388</u>	<u>\$85,399,070</u>
Assets by area		
United States	\$20,221,022	\$20,004,454
Australia/New Zealand	934,368	796,189
Canada	291,760	257,825
Mexico	841,981	681,285
Europe ⁽¹⁾	758,358	466,152
Asia ⁽²⁾	1,796,212	1,948,528
Total consolidated assets	<u>\$24,843,701</u>	<u>\$24,154,433</u>

⁽¹⁾ Europe consists of United Kingdom, Ireland, Germany, Austria, and the Netherlands.

⁽²⁾ Asia consists of Philippines, Malaysia, Singapore, Brunei, and Indonesia.

Reliv' International, Inc. and Subsidiaries

Notes to Consolidated Financial Statements

15. Segment Information (continued)

The Company classifies its sales into three categories of sales products plus handling & freight income. Net sales by product category data for the years ended December 31, 2010 and 2009, follow:

	<u>2010</u>	<u>2009</u>
Net sales by product category		
Nutritional and dietary supplements	\$67,024,233	\$72,556,665
Skin care products	932,631	984,985
Sales aids and other	2,076,716	2,303,949
Handling & freight income	8,714,808	9,553,471
Total net sales	<u>\$78,748,388</u>	<u>\$85,399,070</u>

Consent of Independent Registered Public Accounting Firm

We consent to the incorporation by reference in the following Registration Statements:

1. Registration Statement (Form S-3 No. 333-131974) of Reliv International, Inc., as amended.
2. Registration Statement (Form S-8 No. 333-67921) pertaining to the Reliv International, Inc. 401(k) Plan.
3. Registration Statement (Form S-8 No. 333-107914) pertaining to the Reliv International, Inc. 2003 Stock Option Plan.
4. Registration Statement (Form S-3 No. 333-160374) pertaining to the 2009 Distributor Stock Purchase Plan.
5. Registration Statement (Form S-8 No. 333-170928) pertaining to the Reliv International, Inc. 2009 Incentive Stock Plan.

of our report dated March 17, 2011, with respect to the consolidated financial statements of Reliv International, Inc. and Subsidiaries included in this Annual Report (Form 10-K) of Reliv International, Inc. for the year ended December 31, 2010.

/s/ Ernst & Young LLP

St. Louis, Missouri
March 17, 2011

CERTIFICATION

I, Robert L. Montgomery, certify that:

1. I have reviewed this annual report on Form 10-K of Reliv International, Inc.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - a. Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
 - b. Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
 - c. Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
 - d. Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
5. The registrant's other certifying officer and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - a. All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b. Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: March 17, 2011

/s/ Robert L. Montgomery
Robert L. Montgomery
Chief Executive Officer

CERTIFICATION

I, Steven D. Albright, certify that:

1. I have reviewed this annual report on Form 10-K of Reliv International, Inc.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - a. Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
 - b. Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
 - c. Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
 - d. Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
5. The registrant's other certifying officer and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - a. All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b. Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: March 17, 2011

/s/ Steven D. Albright
Steven D. Albright
Chief Financial Officer

CERTIFICATION PURSUANT TO

18 U.S.C. SECTION 1350.

AS ADOPTED PURSUANT TO

SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

In connection with the Annual Report on Form 10-K of Reliv' International, Inc. (the "Company") for the fiscal year ended December 31, 2010, as filed with the Securities and Exchange Commission on the date hereof (the "Report"), Robert L. Montgomery, as Chief Executive Officer of the Company, and Steven D. Albright, as Chief Financial Officer of the Company, each hereby certifies, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that:

(1) The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and

(2) The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

/s/ Robert L. Montgomery
Robert L. Montgomery
Chief Executive Officer

Date: March 17, 2011

/s/ Steven D. Albright
Steven D. Albright
Chief Financial Officer

Date: March 17, 2011

The foregoing certification is being furnished solely pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, and is not being "filed" as part of the Form 10-K or as a separate disclosure document for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to liability under that section. This certification shall not be deemed to be incorporated by reference into any filing under the Securities Act of 1933, as amended, or the Exchange Act except to the extent that this Exhibit 32 is expressly and specifically incorporated by reference in any such filing.

A signed original of this written statement required by Section 906 has been provided to the Company and will be retained by the Company and furnished to the Securities and Exchange Commission or its staff upon request.

Corporate Headquarters

Reliv International, Inc.
136 Chesterfield Industrial Blvd.
Chesterfield, Missouri 63005
Phone: 636.537.9715
Fax: 636.537.9753

State & Date of Incorporation

Delaware, February 11, 1985

Independent Auditors

Ernst & Young LLP

Fiscal Year-End

December 31

Dividend Reinvestment, Share Purchase & Sale Program

This Program is available to the general public and current shareholders of the Company. If you would like to receive information on this Program, please call American Stock Transfer & Trust Co., toll free, at 888.333.0203.

Stock Exchange Listing

Nasdaq Stock Market® under the symbol RELV.

Annual Meeting

The annual meeting of shareholders will be held at 9:00 a.m. on Thursday, May 26, 2011, at Reliv Corporate Headquarters, 136 Chesterfield Industrial Blvd. Chesterfield, Missouri 63005

Transfer Agent

American Stock Transfer & Trust Co.
59 Maiden Lane, Plaza Level
New York, NY 10038
800.937.5449

Number of Shareholders of Record

1,685 as of March 1, 2011

Shareholder Questions

Communications concerning stock transfer requirements, lost certificates, change of address or dividends should be addressed to American Stock Transfer & Trust Co. at 800.937.5449.

Financial Information

Reliv International maintains a website at www.reliv.com.



Reliv International, Inc.
136 Chesterfield Industrial Blvd.
Chesterfield, Missouri 63005

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